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OM nucleic - nucleic search, using sw model

Run on: June 21, 2005, 02:55:41 ; Search time 546 Seconds
(without alignments)
170.537 Million cell updates/sec

Title: US-10-075-994A-1
Perfect score: 15
Sequence: 1 gtgtccatgatgc 15

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 6054689 seqs, 3103772919 residues

Total number of hits satisfying chosen parameters: 12109378

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database :

Published Applications NA:*

- 1: /cgn2_6/prodata/2/pubpna/US07_PUBCOMB.seq:*
- 2: /cgn2_6/prodata/2/pubpna/PCR_NEM_PUB.seq:*
- 3: /cgn2_6/prodata/2/pubpna/US06_NEM_PUB.seq:*
- 4: /cgn2_6/prodata/2/pubpna/US06_PUBCOMB.seq:*
- 5: /cgn2_6/prodata/2/pubpna/US07_NEM_PUB.seq:*
- 6: /cgn2_6/prodata/2/pubpna/PCRUS_PUBCOMB.seq:*
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- 11: /cgn2_6/prodata/2/pubpna/US09C_PUBCOMB.seq:*
- 12: /cgn2_6/prodata/2/pubpna/US09_NEM_PUB.seq:*
- 13: /cgn2_6/prodata/2/pubpna/US10A_PUBCOMB.seq:*
- 14: /cgn2_6/prodata/2/pubpna/US10B_PUBCOMB.seq:*
- 15: /cgn2_6/prodata/2/pubpna/US10C_PUBCOMB.seq:*
- 16: /cgn2_6/prodata/2/pubpna/US10E_PUBCOMB.seq:*
- 17: /cgn2_6/prodata/2/pubpna/US10F_PUBCOMB.seq:*
- 18: /cgn2_6/prodata/2/pubpna/US10G_PUBCOMB.seq:*
- 19: /cgn2_6/prodata/2/pubpna/US10H_PUBCOMB.seq:*
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- 22: /cgn2_6/prodata/2/pubpna/US11A_PUBCOMB.seq:*
- 23: /cgn2_6/prodata/2/pubpna/US11A_PUBCOMB.seq:*
- 24: /cgn2_6/prodata/2/pubpna/US60_NEM_PUB.seq:*
- 25: /cgn2_6/prodata/2/pubpna/US60_PUBCOMB.seq:*
- 26: /cgn2_6/prodata/2/pubpna/US60_PUBCOMB.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	15	100.0	15	9	US-09-930-283A-1
2	15	100.0	15	10	US-09-930-283A-3
3	15	100.0	15	10	US-09-895-480A-15
4	15	100.0	15	15	US-10-290-545-25
5	15	100.0	15	16	US-10-365-623-16
6	15	100.0	15	17	US-10-075-994A-1
7	15	100.0	15	17	US-10-075-994A-3

8	15	100.0	15	17	US-10-075-994A-4	Sequence 4, Appli
9	15	100.0	15	17	US-10-347-924-1	Sequence 1, Appli
10	15	100.0	15	17	US-10-437-263-25	Sequence 25, Appli
11	15	100.0	15	17	US-10-437-275-25	Sequence 25, Appli
12	15	100.0	15	17	US-10-437-258-25	Sequence 25, Appli
13	15	100.0	15	17	US-10-925-734-15	Sequence 15, Appli
14	15	100.0	20	14	US-10-057-550-49	Sequence 49, Appli
15	15	100.0	20	15	US-10-173-225B-47	Sequence 2, Appli
16	15	100.0	25	9	US-09-930-283A-2	Sequence 2, Appli
17	15	100.0	25	17	US-10-075-994A-2	Sequence 1, Appli
18	15	100.0	25	21	US-10-809-189-113549	Sequence 113549,
19	15	100.0	25	21	US-10-809-189-113550	Sequence 113550,
20	15	100.0	165	9	US-09-728-446-1067	Sequence 1067, Ap
21	15	100.0	478	10	US-09-918-995-4443	Sequence 4443, Ap
22	15	100.0	597	16	US-10-029-386-10799	Sequence 10799, A
23	15	100.0	673	13	US-10-027-632-244006	Sequence 244006,
24	15	100.0	673	13	US-10-027-632-244007	Sequence 244007,
25	15	100.0	673	17	US-10-027-632-244006	Sequence 244006,
26	15	100.0	673	17	US-10-027-632-244007	Sequence 244007,
27	15	100.0	968	19	US-10-767-795-1291	Sequence 1291, Ap
28	15	100.0	1161	19	US-10-437-963-36158	Sequence 36158, A
29	15	100.0	2975	16	US-10-440-341-2	Sequence 2, Appli
30	15	100.0	2977	9	US-09-969-347-207	Sequence 207, App
31	15	100.0	2977	10	US-09-963-131-159	Sequence 159, App
32	15	100.0	2977	14	US-10-057-550-25	Sequence 25, Appli
33	15	100.0	2977	15	US-10-173-225B-64	Sequence 64, Appli
34	15	100.0	2977	16	US-10-371-138-1	Sequence 1, Appli
35	15	100.0	2977	21	US-10-843-641A-8336	Sequence 8336, Ap
36	15	100.0	2977	21	US-10-936-273-29	Sequence 29, Appli
37	15	100.0	3228	21	US-10-926-543-44	Sequence 44, Appli
38	15	100.0	76698	21	US-10-936-273-30	Sequence 30, Appli
39	15	100.0	25	21	US-10-809-189-113548	Sequence 113548
40	14	93.3	223	11	US-09-732-627A-2120	Sequence 2120, Ap
41	14	93.3	323	9	US-10-437-963-4788	Sequence 4788, Ap
42	14	93.3	327	9	US-09-983-965-3423	Sequence 3423, Ap
43	14	93.3	406	20	US-10-425-115-149033	Sequence 149033,
44	14	93.3	475	18	US-10-424-599-103047	Sequence 103047,
45	14	93.3	602	20	US-10-357-930-50238	Sequence 50238, A

ALIGNMENTS

RESULT 1
US-09-930-283A-1
Sequence 1, Application US/09930283A
Patent No. US20020160038A1
GENERAL INFORMATION:
APPLICANT: Kasid, Usha
Gokhale, Prafulla
Dritschilo, Anatoly
Rahman, Agulir
TITLE OF INVENTION: Liposomes containing Oligonucleotides
NUMBER OF SEQUENCES: 3
CORRESPONDENCE ADDRESS:
ADDRESSER: Hendricks and Assoc.
STREET: P.O. Box 2509
CITY: Fairfax
STATE: VA
COUNTRY: US
ZIP: 22031
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/930,283A
FILING DATE: 16-Aug-2001
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 09/354,109
FILING DATE: 1999-07-15

ATTORNEY/AGENT INFORMATION:
NAME: Hendricks, Glenna
REGISTRATION NUMBER: 32,535
REFERENCE/DOCKET NUMBER: Kasid
TELECOMMUNICATION INFORMATION:
TELEPHONE: (703) 591-4470
TELEFAX: (703) 591-4428
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 15 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: unknown
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: YES
SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-09-930-283A-1

Query Match 100.0%; Score 15; DB 9; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTGCTCCATTGATGC 15
Db 1 GTGCTCCATTGATGC 15

RESULT 2
US-09-930-283A-3/C
Sequence 3, Application US/09930283A
Patent No. US20020160038A1
GENERAL INFORMATION:
APPLICANT: Kasid, Usha
Gokhale, Prafulla
Dritschilo, Anatoly
Rahman, Aquilar
TITLE OF INVENTION: Liposomes containing oligonucleotides
NUMBER OF SEQUENCES: 3
CORRESPONDENCE ADDRESS:
ADDRESSEE: Hendricks and Assoc.
STREET: P.O. Box 2509
CITY: Fairfax
STATE: VA
COUNTRY: US
ZIP: 22031
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/930,283A
FILING DATE: 16-Aug-2001
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 09/354,109
FILING DATE: 1998-07-15
ATTORNEY/AGENT INFORMATION:
NAME: Hendricks, Glenna
REGISTRATION NUMBER: 32,535
REFERENCE/DOCKET NUMBER: Kasid
TELECOMMUNICATION INFORMATION:
TELEPHONE: (703) 591-4470
TELEFAX: (703) 591-4428
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 15 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: unknown
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO

ANTI-SENSE: YES
SEQUENCE DESCRIPTION: SEQ ID NO: 3:
US-09-930-283A-3

Query Match 100.0%; Score 15; DB 9; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTGCTCCATTGATGC 15
Db 15 GTGCTCCATTGATGC 1

RESULT 3
US-09-895-480A-15
Sequence 15, Application US/09895480A
Publication NO. US20030129221A1
GENERAL INFORMATION:
APPLICANT: Inex Pharmaceuticals Inc.
TITLE OF INVENTION: High Efficiency Encapsulation of Charged Therapeutic Agents in Lipid Vesicles
NUMBER OF SEQUENCES: 17
CORRESPONDENCE ADDRESS:
ADDRESSEE: Oppedahl & Larson LLP
STREET: PO Box 5068
CITY: Dillon
STATE: CO
COUNTRY: US
ZIP: 80435
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.5 inch, 1.44 MB
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS 5.0
SOFTWARE: Word Perfect
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/895,480A
FILING DATE: 29-Jun-2001
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: <Unknown>
FILING DATE: <Unknown>
ATTORNEY/AGENT INFORMATION:
NAME: <Unknown>
REGISTRATION NUMBER: <Unknown>
REFERENCE/DOCKET NUMBER: <Unknown>
TELECOMMUNICATION INFORMATION:
TELEPHONE: <Unknown>
TELEFAX: <Unknown>
INFORMATION FOR SEQ ID NO: 15:
SEQUENCE CHARACTERISTICS:
LENGTH: 15
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
HYPOTHETICAL: no
ANTI-SENSE: yes
SEQUENCE DESCRIPTION: SEQ ID NO: 15:
US-09-895-480A-15

Query Match 100.0%; Score 15; DB 10; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTGCTCCATTGATGC 15
Db 1 GTGCTCCATTGATGC 15

RESULT 4
US-10-290-545-25

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Sequence 25, Application US/10290545
; Publication No. US20030125292A1
; GENERAL INFORMATION:
; APPLICANT: Semple, Sean
; APPLICANT: Klimuk, Sandy
; APPLICANT: Yuan, Zuan-Ning
; TITLE OF INVENTION: Improved Mucosal Vaccines and Methods for Using the Same
; FILE REFERENCE: A-71854/TRL/XG
; CURRENT APPLICATION NUMBER: US/10/290,545
; CURRENT FILING DATE: 2002-11-07
; NUMBER OF SEQ ID NOS: 30
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 25
; LENGTH: 15
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-290-545-25

Query Match          100.0%; Score 15; DB 15; Length 15;
Best Local Similarity 100.0%; Pred. No. 1,3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 GTGCTCCATTGATGC 15
DB      1 GTGCTCCATTGATGC 15

RESULT 5
US-10-365-623-16
; Sequence 16, Application US/10365623
; Publication No. US20030166512A1
; GENERAL INFORMATION:
; APPLICANT: Xie, Dong
; TITLE OF INVENTION: Protein Carrier System for Therapeutic Oligonucleotides
; FILE REFERENCE: 63024,000001
; CURRENT APPLICATION NUMBER: US/10/365,623
; CURRENT FILING DATE: 2003-02-13
; NUMBER OF SEQ ID NOS: 23
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 16
; LENGTH: 15
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Antisense oligonucleotide directed against human c-rac-1 protein
US-10-365-623-16

Query Match          100.0%; Score 15; DB 16; Length 15;
Best Local Similarity 100.0%; Pred. No. 1,3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 GTGCTCCATTGATGC 15
DB      1 GTGCTCCATTGATGC 15

RESULT 6
US-10-075-994A-1
; Sequence 1, Application US/10075994A
; Publication No. US20030215489A1
; GENERAL INFORMATION:
; APPLICANT: KASID, Usha
; APPLICANT: GOKHALE, Prafulla
; APPLICANT: PIE, Jin
; APPLICANT: MEWANI, Rajshree
; APPLICANT: AHMAD, Imran
; APPLICANT: DRITSCHILLO, Anatoly
; APPLICANT: RAHMAN, Aquilur
; TITLE OF INVENTION: CHEMOSENSITIZING WITH LIPOSOMES CONTAINING OLIGONUCLEOTIDES
; FILE REFERENCE: 219604
; CURRENT APPLICATION NUMBER: US/10/075,994A
; CURRENT FILING DATE: 2002-02-15
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NUMBER OF SEQ ID NOS: 7
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 1
; LENGTH: 15
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic
US-10-075-994A-1

Query Match          100.0%; Score 15; DB 17; Length 15;
Best Local Similarity 100.0%; Pred. No. 1,3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 GTGCTCCATTGATGC 15
DB      1 GTGCTCCATTGATGC 15

RESULT 7
US-10-075-994A-3/c
; Sequence 3, Application US/10075994A
; Publication No. US20030215489A1
; GENERAL INFORMATION:
; APPLICANT: KASID, Usha
; APPLICANT: GOKHALE, Prafulla
; APPLICANT: PIE, Jin
; APPLICANT: MEWANI, Rajshree
; APPLICANT: AHMAD, Imran
; APPLICANT: DRITSCHILLO, Anatoly
; APPLICANT: RAHMAN, Aquilur
; TITLE OF INVENTION: CHEMOSENSITIZING WITH LIPOSOMES CONTAINING OLIGONUCLEOTIDES
; FILE REFERENCE: 219604
; CURRENT APPLICATION NUMBER: US/10/075,994A
; CURRENT FILING DATE: 2002-02-15
; NUMBER OF SEQ ID NOS: 7
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 3
; LENGTH: 15
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic
US-10-075-994A-3

Query Match          100.0%; Score 15; DB 17; Length 15;
Best Local Similarity 100.0%; Pred. No. 1,3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 GTGCTCCATTGATGC 15
DB      15 GTGCTCCATTGATGC 1

RESULT 8
US-10-075-994A-4
; Sequence 4, Application US/10075994A
; Publication No. US20030215489A1
; GENERAL INFORMATION:
; APPLICANT: KASID, Usha
; APPLICANT: GOKHALE, Prafulla
; APPLICANT: PIE, Jin
; APPLICANT: MEWANI, Rajshree
; APPLICANT: AHMAD, Imran
; APPLICANT: DRITSCHILLO, Anatoly
; APPLICANT: RAHMAN, Aquilur
; TITLE OF INVENTION: CHEMOSENSITIZING WITH LIPOSOMES CONTAINING OLIGONUCLEOTIDES
; FILE REFERENCE: 219604
; CURRENT APPLICATION NUMBER: US/10/075,994A
; CURRENT FILING DATE: 2002-02-15
; NUMBER OF SEQ ID NOS: 7
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 4
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LENGTH: 15
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Synthetic
US-10-075-994A-4

Query Match 100.0%; Score 15; DB 17; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTGCTCCATTGATGC 15
DB 1 GTGCTCCATTGATGC 15

RESULT 9

US-10-347-924-1
Sequence 1, Application US/10347924
Publication No. US20030229040A1
GENERAL INFORMATION:

APPLICANT: Kasid, Usha
APPLICANT: Gokhale, Prafulla
APPLICANT: Zhang, Chuabo
APPLICANT: Dristschilo, Anatoly
APPLICANT: Rahman, Aquilur
TITLE OF INVENTION: CATIONIC LIPOSOMAL DELIVERY SYSTEM AND THERAPEUTIC USE THEREOF
FILE REFERENCE: 220807
CURRENT APPLICATION NUMBER: US/10/347,924
CURRENT FILING DATE: 2003-01-21
PRIOR APPLICATION NUMBER: US 09/354,109
PRIOR FILING DATE: 1999-07-15
PRIOR APPLICATION NUMBER: US 08/957,327
PRIOR FILING DATE: 1997-10-24
PRIOR APPLICATION NUMBER: US 60/041,192
PRIOR FILING DATE: 1997-03-21
NUMBER OF SEQ ID NOS: 1
SOFTWARE: PatentIn version 3.2
SEQ ID NO 1
LENGTH: 15
TYPE: DNA
ORGANISM: Artificial
FEATURE:
OTHER INFORMATION: Oligonucleotide
US-10-347-924-1

Query Match 100.0%; Score 15; DB 17; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTGCTCCATTGATGC 15
DB 1 GTGCTCCATTGATGC 15

RESULT 10

US-10-437-263-25
Sequence 25, Application US/10437263
Publication No. US20040009943A1
GENERAL INFORMATION:

APPLICANT: Semple, Sean
APPLICANT: Tam, Ying K.
APPLICANT: Chikh, Ghania
APPLICANT: Hope, Michael J.
TITLE OF INVENTION: PATHOGEN VACCINES AND METHODS FOR USING THE SAME
FILE REFERENCE: A-72216/TAL
CURRENT APPLICATION NUMBER: US/10/437,263
CURRENT FILING DATE: 2003-05-12
PRIOR APPLICATION NUMBER: 60/379,343
PRIOR FILING DATE: 2002-05-10
PRIOR APPLICATION NUMBER: 60/460,646
PRIOR FILING DATE: 2003-04-04
PRIOR APPLICATION NUMBER: 60/454,298

PRIOR FILING DATE: 2003-03-12
NUMBER OF SEQ ID NOS: 34
SOFTWARE: PatentIn version 3.2
SEQ ID NO 25
LENGTH: 15
TYPE: DNA
ORGANISM: Homo sapiens
US-10-437-263-25

Query Match 100.0%; Score 15; DB 17; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTGCTCCATTGATGC 15
DB 1 GTGCTCCATTGATGC 15

RESULT 11

US-10-437-275-25
Sequence 25, Application US/10437275
Publication No. US20040009944A1
GENERAL INFORMATION:

APPLICANT: Tam, Ying K.
APPLICANT: Semple, Sean
APPLICANT: Klimuk, Sandra
APPLICANT: Chikh, Ghania
TITLE OF INVENTION: METHYLATED IMMUNOSTIMULATORY OLIGONUCLEOTIDES AND METHODS OF
FILE REFERENCE: A-72158/TAL
CURRENT APPLICATION NUMBER: US/10/437,275
CURRENT FILING DATE: 2003-05-12
PRIOR APPLICATION NUMBER: 60/379,343
PRIOR FILING DATE: 2002-05-10
PRIOR APPLICATION NUMBER: 60/460,646
PRIOR FILING DATE: 2003-04-04
PRIOR APPLICATION NUMBER: 10/290,545
PRIOR FILING DATE: 2002-11-07
NUMBER OF SEQ ID NOS: 32
SOFTWARE: PatentIn version 3.2
SEQ ID NO 25
LENGTH: 15
TYPE: DNA
ORGANISM: Homo sapiens
US-10-437-275-25

Query Match 100.0%; Score 15; DB 17; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTGCTCCATTGATGC 15
DB 1 GTGCTCCATTGATGC 15

RESULT 12

US-10-437-258-25
Sequence 25, Application US/10437258
Publication No. US20040013649A1
GENERAL INFORMATION:

APPLICANT: Semple, Sean
APPLICANT: Tam, Ying K.
APPLICANT: Klimuk, Sandra
APPLICANT: Chikh, Ghania
TITLE OF INVENTION: CANCER VACCINES AND METHODS OF USING THE SAME
FILE REFERENCE: A-72252/TAL
CURRENT APPLICATION NUMBER: US/10/437,258
CURRENT FILING DATE: 2003-05-12
PRIOR APPLICATION NUMBER: 60/379,343
PRIOR FILING DATE: 2002-05-10
PRIOR APPLICATION NUMBER: 60/460,646
PRIOR FILING DATE: 2003-04-04
PRIOR APPLICATION NUMBER: 60/454,298

PRIOR FILING DATE: 2003-03-12
NUMBER OF SEQ ID NOS: 34
SOFTWARE: PatentIn version 3.2
SEQ ID NO: 25
LENGTH: 15
TYPE: DNA
ORGANISM: Homo sapiens
US-10-437-258-25

Query Match 100.0%; Score 15; DB 17; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 GTGCTCCATTGATGC 15
Db 1 GTGCTCCATTGATGC 15

RESULT 13
US-10-925-734-15
Sequence 15, Application US/10925734
Publication No. US2005000869A1
GENERAL INFORMATION:
APPLICANT: Inex Pharmaceuticals Inc.
TITLE OF INVENTION: High Efficiency Encapsulation of Charged
Therapeutic
Agents in
Lipid Vesicles

NUMBER OF SEQUENCES: 17
CORRESPONDENCE ADDRESS:
ADDRESSER: Oppedahl & Larson LLP
STREET: PO Box 5068
CITY: Dillon
STATE: CO
COUNTRY: US
ZIP: 80435

COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.5 inch, 1.44 Mb

COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS 5.0
SOFTWARE: Word Perfect

CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/10/925,734
FILING DATE: 24-Aug-2004
CLASSIFICATION: <Unknown>

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/09/895,480
FILING DATE: 29-Jun-2001

ATTORNEY/AGENT INFORMATION:
NAME: <Unknown>

REGISTRATION NUMBER: <Unknown>
REFERENCE/DOCKET NUMBER: <Unknown>
TELECOMMUNICATION INFORMATION:
TELEPHONE: <Unknown>
TELEFAX: <Unknown>
TELEX: <Unknown>

INFORMATION FOR SEQ ID NO: 15:
SEQUENCE CHARACTERISTICS:
LENGTH: 15
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
HYPOTHEICAL: no
ANTI-SENSE: yes

SEQUENCE DESCRIPTION: SEQ ID NO: 15:
US-10-925-734-15

Query Match 100.0%; Score 15; DB 21; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 GTGCTCCATTGATGC 15

Db 1 GTGCTCCATTGATGC 15

RESULT 14
US-10-057-550-49
Sequence 49, Application US/10057550
Publication No. US20030032607A1
GENERAL INFORMATION:
APPLICANT: Monia, Brett P.
TITLE OF INVENTION: Antisense Oligonucleotide Modulation of raf Gene Expression
FILE REFERENCE:
CURRENT APPLICATION NUMBER: US/10/057,550
CURRENT FILING DATE: 2002-01-25
PRIOR APPLICATION NUMBER: 09/506,073
PRIOR FILING DATE: 2000-02-18
PRIOR APPLICATION NUMBER: US 09/143,214
PRIOR FILING DATE: 1998-08-28
PRIOR APPLICATION NUMBER: PCT/US98/13961
PRIOR FILING DATE: 1998-07-06
PRIOR APPLICATION NUMBER: US 08/888,982
PRIOR FILING DATE: 1997-07-07
PRIOR APPLICATION NUMBER: US 08/756,806
PRIOR FILING DATE: 1996-11-26
PRIOR APPLICATION NUMBER: PCT/US95/07111
PRIOR FILING DATE: 1995-05-31
PRIOR APPLICATION NUMBER: US 08/250,856
PRIOR FILING DATE: 1994-05-31
NUMBER OF SEQ ID NOS: 130
SEQ ID NO: 49
LENGTH: 20
TYPE: DNA
ORGANISM: artificial sequence
FEATURE:
OTHER INFORMATION: antisense sequence
US-10-057-550-49

Query Match 100.0%; Score 15; DB 14; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 GTGCTCCATTGATGC 15
Db 5 GTGCTCCATTGATGC 19

RESULT 15
US-10-173-225B-47
Sequence 47, Application US/10173225B
Publication No. US20030119769A1
GENERAL INFORMATION:
APPLICANT: Monia, Brett P.
TITLE OF INVENTION: Antisense Oligonucleotide Modulation of raf Gene Expression
FILE REFERENCE: ISPH-0665
CURRENT APPLICATION NUMBER: US/10/173,225B
CURRENT FILING DATE: 2002-12-06
PRIOR APPLICATION NUMBER: US 10/057,550
PRIOR FILING DATE: 2002-01-25
PRIOR APPLICATION NUMBER: US 09/143,214
PRIOR FILING DATE: 1998-08-28
PRIOR APPLICATION NUMBER: PCT/US98/13961
PRIOR FILING DATE: 1998-07-06
PRIOR APPLICATION NUMBER: US 08/888,982
PRIOR FILING DATE: 1997-07-07
PRIOR APPLICATION NUMBER: US 08/756,806
PRIOR FILING DATE: 1996-11-26
PRIOR APPLICATION NUMBER: PCT/US95/07111
PRIOR FILING DATE: 1995-05-31
PRIOR APPLICATION NUMBER: US 08/250,856
PRIOR FILING DATE: 1994-05-31
NUMBER OF SEQ ID NOS: 109
SEQ ID NO: 47
LENGTH: 20

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OM nucleic - nucleic search, using sw model

Run on: June 21, 2005, 03:49:31 ; Search time 427 Seconds
(without alignments)
207.953 Million cell updates/sec

Title: US-10-075-994A-1
Perfect score: 15
Sequence: 1 gtgtccatcgcgc 15

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues
Total number of hits satisfying chosen parameters: 4167226

Minimum DB seq length: 0
Maximum DB seq length: 50

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
1: N Geneseq_16Dec04:*
2: geneseqn1980s:*
3: geneseqn1990s:*
4: geneseqn2000s:*
5: geneseqn2001as:*
6: geneseqn2001bs:*
7: geneseqn2002as:*
8: geneseqn2002bs:*
9: geneseqn2003as:*
10: geneseqn2003bs:*
11: geneseqn2003ds:*
12: geneseqn2004as:*
13: geneseqn2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	15	100.0	15	2	AAV54043 Human ant
2	15	100.0	15	2	AAV99435 Antisense
3	15	100.0	15	3	AAZ98661 Human C-r
4	15	100.0	15	6	AAZ22797 Human C-r
5	15	100.0	15	9	ACC58517 Oligonuc
6	15	100.0	15	9	ADA24233 Human C-r
7	15	100.0	15	10	ADB97458 Sense (AT
8	15	100.0	15	10	ADB97456 Antisense
9	15	100.0	15	10	ADF82830 Immunost
10	15	100.0	15	12	ADG90171 Human C-r
11	15	100.0	15	12	ADG39690 Oligonuc
12	15	100.0	15	12	ADF32025 Antisense
13	15	100.0	15	12	ADF42926 Methylate
14	15	100.0	15	15	ADL70154 Oligonuc
15	15	100.0	15	13	ADR88950 Anti C-ra
16	15	100.0	20	2	AAZ7527 Mouse/rat
17	15	100.0	20	2	AAZ11557 Mouse and
18	15	100.0	20	3	AAZ73535 Mouse and
19	15	100.0	20	6	AAZ44760 Mouse/rat
20	15	100.0	20	10	ADP09751 Mouse/rat

21	15	100.0	20	10	ACD42120 Antisense
22	15	100.0	25	10	ADB97457 Oligo use
23	13	86.7	17	2	AAV90935 Human C-r
24	13	86.7	20	2	AAZ7482 Human C-r
25	13	86.7	20	2	AAZ59716 Human C-r
26	13	86.7	20	2	AAZ62145 Human C-r
27	13	86.7	20	2	AAZ11512 Human C-r
28	13	86.7	20	3	AAZ73490 Human C-r
29	13	86.7	20	6	AAZ44715 Human C-r
30	13	86.7	20	10	ADP09706 Human C-r
31	13	86.7	20	10	ACD42073 Antisense
32	12.4	82.7	20	2	AAZ16555 Position
33	12.4	82.7	20	2	AAZ16583 Position
34	12.4	82.7	20	2	AAZ23613 Homo sapi
35	12.4	82.7	20	6	AB194231 Capture o
36	12.4	82.7	24	6	AB185032 Capture o
37	12.4	82.7	24	6	AB185033 Capture o
38	12.4	82.7	27	2	AAV93971 Human IL-
39	12	80.0	20	2	AAZ27483 Human C-r
40	12	80.0	20	2	AAZ86617 Rat C-raf
41	12	80.0	20	3	AAZ11513 Human C-r
42	12	80.0	20	3	AAZ92027 C-raf tar
43	12	80.0	20	3	AAZ92026 C-raf tar
44	12	80.0	20	3	AAZ73491 Human C-r
45	12	80.0	20	6	AAZ44716 Human C-r

ALIGNMENTS

RESULT 1
AAV54043
ID AAV54043 standard; DNA; 15 BP.
XX
AC AAV54043;
XX
DT 02-DEC-1998 (first entry)
XX
DE Human antisense c-raf-1 oligodeoxyribonucleotide.
XX
KW Human; antisense; c-raf-1; oligodeoxyribonucleotide; ODN/oligo;
RV tumour tissue; cancer; radiation therapy; radiosensitise; antisense;
KW liposome carrier system; ss.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT modified_base 1
FT /note= "N-terminal base is phosphothioated"
FT /tag= a
FT modified_base 15
FT /note= "C-terminal base is phosphothioated"
FT /tag= b
XX W09843095-A1.
XX PD 01-OCT-1998.
XX PF 19-MAR-1998; 98WO-US005303.
XX PR 21-MAR-1997; 97US-0041192P.
XX PR 24-OCT-1997; 97US-00957327.
XX PA (GEO) UNIV GORGETOWN.
XX PI Kasid U, Gokhale P, Dritschilo A, Rahman A;
XX WPI; 1998-532155/45.
XX PT New cationic liposome composition containing raf oligodeoxynucleotide -
XX PT can be used to directly target tumour tissue and is useful in the
XX PT radiation therapy of cancers.

PS Claim 4; Page 21; 25pp; English.

XX This is the nucleotide sequence of the human antisense c-raf-1

CC oligodeoxynucleotide (ODN/oligo), used in the method of the invention

CC to directly target tumour tissue, and in cancer radiation therapy. The

CC products can be used in a method of radiosensitising tumour tissue by

CC addition of an antisense oligonucleotide of maximum 40 bases containing

CC ODN/oligo. The liposome carrier system directly targets tumour tissue and

CC has the potential for use in the radiation therapy of cancers

XX

SQ Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;

QY Query Match 100.0%; Score 15; DB 2; Length 15;

Best Local Similarity 100.0%; Pred. No. 1.1e+02;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 GTGCTCCATTGATGC 15

RESULT 2

AAV99435

ID AAV99435 standard; DNA; 15 BP.

AC AAV99435;

XX

DT 22-MAR-1999 (first entry)

XX

DE Antisense oligonucleotide directed against c-raf-1 protein kinase gene.

XX

KM Antisense oligonucleotide; human c-raf-1 protein kinase gene;

KM phosphorothioate; phosphodiester; lipid-encapsulation; tumour;

KM aberrant gene expression; treatment; inflammation; infection; ss.

XX

OS Synthetic.

OS Homo sapiens.

XX

XX

XX Key Location/Qualifiers

FT modified_base 1..15

FT /+tag= a

FT /note= "phosphorothioate or phosphodiester bonds"

XX

PN WO9851278-A2.

XX

PD 19-NOV-1998.

XX

PF 14-MAY-1998; 98WO-CA000485.

XX

PR 14-MAY-1997; 97US-00856374.

XX

XX (INEX-) INEX PHARM CORP.

PA

PI Sempje SC, Klimuk SK, Harasym T, Hope MJ, Ansell SM, Cullis P,

PI Scherrer P, Debeyer D;

XX

DR WPI; 1999-045179/04.

XX

PT Composition containing lipid-encapsulated therapeutic agent - useful,

PT e.g. for delivering antisense molecules or ribozymes or treating diseases

PT associated with aberrant gene expression.

XX

PS Disclosure; Page 23; 99pp; English.

XX

XX The present sequence represents an antisense oligonucleotide directed

CC against the human c-raf-1 protein kinase gene. The oligonucleotide can

CC have either phosphorothioate or phosphodiester bonds. The oligonucleotide

CC is lipid-encapsulated using the method of the invention. A composition

CC comprising lipid-encapsulated particles of a therapeutic agent, e.g.

CC antisense oligonucleotides, is prepared by mixing at least 2 lipids with

CC buffered aqueous solution of charged therapeutic agent to form an

CC intermediate mixture of lipid-encapsulated particles, and changing the pH

CC of the mixture to neutralise at least some of the external surface

CC charges on the particles. One lipid has a (de)protonatable group with Ka

CC such that the lipid is charged at a first pH but neutral at a second pH

CC (particularly near physiological pH) and the buffer maintains this lipid

CC in the charged form (i.e. cationic when the therapeutic agent is anionic

CC in the buffer, or vice versa). The second lipid prevents particle

CC aggregation during formation of the lipid-therapeutic agent particles.

CC The composition is used to introduce therapeutic agents into cells, in

CC vivo or in vitro, particularly to treat or prevent diseases associated

CC with aberrant gene expression in mammals, specifically tumours,

CC inflammation or infection

XX

SQ Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;

QY Query Match 100.0%; Score 15; DB 2; Length 15;

Best Local Similarity 100.0%; Pred. No. 1.1e+02;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 GTGCTCCATTGATGC 15

RESULT 3

AAZ98661

ID AAZ98661 standard; DNA; 15 BP.

AC AAZ98661;

XX

DT 05-JUN-2000 (first entry)

XX

DE Human c-raf-1 PK therapeutic antisense oligonucleotide sequence ATG-AS.

XX

KM Antisense oligonucleotide; phosphorothioate; inflammatory disease;

KM tumour; gene therapy; aberrant gene expression; treatment;

KM infectious disease; protein kinase C alpha; c-raf-1 protein kinase; ss.

XX

OS Homo sapiens.

XX

XX

XX Key Location/Qualifiers

FT misc_feature 1..15

FT /+tag= a

FT /note= "Optionally phosphorothioate internucleotide linkages"

XX

PN CA2271582-A1.

XX

PD 14-NOV-1999.

XX

PF 13-MAY-1999; 99CA-02271582.

XX

PR 14-MAY-1998; 98US-00078955.

XX

XX (KLIM/) KLIMUK S K.

PA (HARA/) HARASYM T.

PA (HOPE/) HOPE M J.

PA (ANSEL/) ANSELL S M.

PA (CULL/) CULLIS P R.

PA (MOKW/) MOK W K.

PA (SCHE/) SCHERRER P.

PA (SEMP/) SEMPLE S C.

XX

PI Klimuk SK, Harasym T, Hope MJ, Ansell SM, Cullis PR, Mok WWK;

PI Scherrer P, Sempje SC;

XX

DR WPI; 2000-225058/20.

XX

PT A method for delivering antisense oligonucleotides to cells using lipid

PT capsules comprising steric barrier lipids.

XX

PS Example 5; Page 57; 99pp; English.

XX

CC This sequence represents an antisense oligonucleotide sequence which has

CC human c-raf-1 protein kinase as its target gene. The oligonucleotide is

used in a method for delivering lipid encapsulated therapeutic agents (i.e. antisense oligonucleotides) to mammals. The lipid capsule comprises steric barrier lipids that prevent particle aggregation during lipid nucleic acid formation. The method may be used for the delivery of therapeutic agents to mammalian cells, it is especially suitable for delivering nucleic acid molecules, and in particular antisense molecules which may be administered to down regulate the expression of aberrant genes. The aberrant gene may be ICAM-1, c-myc, c-mycb, ras, raf, erb-B-2, PKC-alpha, IGF-1R, EGFR, VEGF and/or VEGF-R-1. The method may be used for the treatment of tumours, inflammatory diseases and/or infectious diseases

Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 15; DB 3; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 GTGCTCCATTGATGC 15
1 GTGCTCCATTGATGC 15

RESULT 4
AAD22797 standard; DNA; 15 BP.

AAD22797;
26-FEB-2002 (first entry)

Human c-raf-1 protein kinase antisense oligonucleotide, ATG-AS.

Treatment; tumour; lipid-therapeutic agent particle; sphingomyelin; distearylphosphatidylcholine; palmitoylcholeoyl phosphatidylcholine; DSPC; POPC; 1,2-dioleoyl-sn-3-phosphoethanolamine; cholesterol; SM; DOPC; inflammation; c-raf-1 protein kinase gene; human; infectious disease; ss.

Homo sapiens.

Key Location/Qualifiers
modified_base 1..20
/*tag= a
/mod_base= OTHER
/note= "Optionally phosphorothioate backbone"

US6287591-B1.
11-SEP-2001.
14-MAY-1998; 98US-00078954.
14-MAY-1997; 97US-00856374.
(INEX-) INEX PHARM CORP.
Sempke SC, Klimuk SK, Haraaym T, Hope MJ, Ansell SM, Cullis P; Scherrer P, Debeyre D;
WPI; 2002-024658/03.
Composition useful for treatment of e.g. tumors comprises particles comprising lipid portion and a charged therapeutic agent.
Disclosure; Col 15-16; 48pp; English.

The invention relates to a composition useful for treatment of e.g. tumours. The composition comprises lipid-therapeutic agent particles comprising a lipid portion and a charged therapeutic agent which is encapsulated in the lipid portion. The lipid portion comprises a first lipid component selected from lipids containing a protonatable or deprotonatable (preferably protonatable) group that has a pKa such that the lipid is in charged form at a first pH and in neutral form at a

second pH. The pKa of lipid component is from 4-11. The first lipid component is further selected such that the charged form is cationic when the therapeutic agent is anionic and vice versa; the second lipid component is selected from lipids that prevent particle aggregation during lipid-therapeutic agent particles formation and which exchange out the lipid particle at a rate greater than PEG-Cerc20; third lipid component is a neutral lipid selected from distearylphosphatidylcholine (DSPC), palmitoylcholeoyl phosphatidylcholine (POPC), 1,2-dioleoyl-sn-3-phosphoethanolamine (DOPC) or SM (sphingomyelin) and a fourth lipid component which is cholesterol. Compositions of the invention are used for treatment or prevention of a disease caused by aberrant expression of a gene preferably ICAM-1 (intracellular adhesion molecule-1), c-myc, c-mycb, ras, raf, erb-B-2, PKC-alpha (phosphokinase C-alpha), IGF-1R (insulin growth factor 1-receptor), bcl-2, EGFR (epidermal growth factor receptor), VEGF and VEGF-R-1 (vascular endothelial growth factor receptor 1) in a mammal or by inflammations such as tumour or an infectious disease. The present sequence is an antisense oligonucleotide targeted to human c-raf-1 protein kinase gene

Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 15; DB 6; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 GTGCTCCATTGATGC 15
1 GTGCTCCATTGATGC 15

RESULT 5
ACC58517 standard; DNA; 15 BP.

ACC58517;
26-AUG-2003 (first entry)

Oligonucleotide ODN #25 (hc-Raf1).

Lipid nucleic acid; LNA; mucosal; vaccine; immunostimulant; human; C-Raf-s; ss.

Homo sapiens.

Key Location/Qualifiers
modified_base 1..15
/*tag= a
/mod_base= OTHER
/note= "OTHER= optional phosphorothioate nucleotides"

WO2003039595-A2.
15-MAY-2003.
07-NOV-2002; 2002WO-CA001717.
07-NOV-2001; 2001US-0337522P.
10-MAY-2002; 2002US-0379343P.
(INEX-) INEX PHARM CORP.
Sempke S, Klimuk S, Yuan Z;
WPI; 2003-493235/46.
Improved mucosal adjuvant useful in the preparation of vaccine for stimulating an immune response comprises a lipid-nucleic acid formulation containing a nucleic acid component encapsulated by a lipid.
Disclosure; Page 21; 71pp; English.

The present sequence is that of oligodeoxynucleotide ODN #25 (hc-Raf-1)

CC for human C-Raf-s. It is an example of an ODN that can be used in lipid-
CC nucleic acid (LNA) formulations of the invention comprising a lipid
CC component and a nucleic acid component. The invention is based on the
CC discovery that such LNA formulations associated with a target antigen
CC stimulate enhanced mucosal immune responses, especially Iga production,
CC directed to that target antigen in vivo as compared to the target antigen
CC alone or mixed with free or unencapsulated forms of the ODN. Claimed
CC improved mucosal vaccines comprise an LNA formulation with at least one
CC antigen, the LNA formulation comprising a lipid component that
CC encapsulates the nucleic acid component, with the lipid and nucleic acid
CC components acting synergistically to stimulate antigen-specific IgG
CC production in a mammal

SQ Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 15; DB 9; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GTGCTCCATTGATGC 15
|||
1 GTGCTCCATTGATGC 15

Db 1 GTGCTCCATTGATGC 15

RESULT 6
ADA24233 standard; DNA; 15 BP.

AC ADA24233;

DT 20-NOV-2003 (first entry)

DE Human c-raf-1 protein kinase antisense oligonucleotide SEQ ID NO:16.

KM therapeutic oligonucleotide; double-stranded RNA; dsRNA; mobile protein;
KM cytosolic; immunosuppressive; virucide; anti-HIV; antibacterial;
KM cardiac; hyperproliferation; cancer; haematological; metastatic;
KM autoimmune disease; infection; endocrine; neural; cardiovascular;
KM pulmonary; reproductive system disorder; endocytosis; metabolic process;
KM murine; intracellular adhesion molecule 1; ICM-1;
KM antisense oligonucleotide; phosphorothioate; ss.

OS Synthetic.
XX Homo sapiens.

XX Key Location/Qualifiers

FT modified_base 1..15
FT /*tag= a
FT /mod_base= OTHER
FT /note= "optionally phosphorothioate backbone"

PN WO2003069306-A2.

PD 21-AUG-2003.

PF 13-FEB-2003; 2003MO-US004323.

PR 13-FEB-2002; 2002US-0356053P.

PA (MEDB-) MEDBRIDGE INC.

PI Xie D;

DR WPI; 2003-646491/61.

PT Treating diseases with oligonucleotides or interfering RNA, useful e.g.
PT for cancer or autoimmune diseases, covalently coupled to mobile proteins,
PT in vivo or in vitro.

PS Claim 128; Page 12; 42pp; English.

CC The present invention describes a method for treating a disease by
CC administering: (a) a therapeutic oligonucleotide (TON) or double-stranded

CC RNA (dsRNA) that includes a reactive group (RG) that can react with a
CC mobile protein (MP) to form a covalent conjugate of TON/dsRNA and MP; or
CC (b) TON or dsRNA already conjugated to MP through a covalent bond. Also
CC described: (1) TON of 15-30 bases that includes (i) a part that binds to
CC target RNA or DNA and (ii) RG; (2) TON or 15-30 bases that includes a
CC part that binds to target RNA or DNA and is conjugated to MP through a
CC covalent link; (3) dsRNA that includes RG; and (4) dsRNA that is
CC conjugated to MP through a covalent link. TON have cytosolic,
CC immunosuppressive, virocidic, anti-HIV, antibacterial and cardiant
CC activities. The method is used to treat, or prevent, hyperproliferation
CC (particularly cancers, solid or haematological, including prevention of
CC metastatic spread); autoimmune diseases; viral or bacterial infections;
CC endocrine, neural, cardiovascular, pulmonary or reproductive system
CC disorders. Also where TON or dsRNA are labelled, they can be used for
CC diagnosis and monitoring of therapy. When linked to a mobile protein,
CC TON/dsRNA have better cell entry (via endocytosis or other parts of the
CC mobile protein metabolic process) and longer therapeutic life, increased
CC from hours to weeks (the result of increased resistance to nucleases),
CC without loss of affinity for the target. In many cases immune response to
CC TON/dsRNA is also reduced, as is non-specific binding to endogenous
CC proteins. The present sequence represents a human c-raf-1 antisense
CC oligonucleotide, which is a specifically claimed TON from the present
CC invention.

SQ Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 15; DB 9; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GTGCTCCATTGATGC 15
|||
1 GTGCTCCATTGATGC 15

Db 1 GTGCTCCATTGATGC 15

RESULT 7

ADB97458/c standard; DNA; 15 BP.

AC ADB97458;

DT 04-DEC-2003 (first entry)

DE Sense (ATG-S) raf ODN oligodeoxyribonucleotide.

KM antisense; ATG-S; raf ODN; chemosensitisation; tumour tissue;

KM chemotherapeutic agent; cationic liposome; cationic lipid;

KM phosphatidylcholine; cholesterol; liposome;

KM dimethyldioctadecyl ammonium bromide; DDAB;

KM dimyristoyl trimethyl ammonium propane; DM7AP; phosphatidylcholine; PC;

KM cholesterol; cancer; leukaemia; lymphoma; myeloma; carcinoma; sarcoma;

KM combination therapy; pre-cancerous lesion; chemotherapy; ss.

OS Unidentified.

PN WO2003070221-A1.

PD 28-AUG-2003.

PF 14-FEB-2003; 2003MO-US004681.

PR 15-FEB-2002; 2002US-00075994.

PA (GEOU) UNIV GEORGETOWN.

PI (NEOP-) NEOPHARM INC.

PT Kasid U, Gokhale P, Pel J, Newani R, Ahmad I, Drischilo A;
PT Rahman A;
WPI; 2003-689738/65.

CC Chemosensitization of tumor tissue, useful for treating cancer, e.g.
CC leukemia, lymphoma or myeloma, comprises administering a chemotherapeutic

PT agent and cationic liposomes containing oligonucleotides.
 XX
 PS Example 1; Page 18; 77pp; English.
 XX
 CC The invention relates to a novel method for the chemosensitisation of
 CC tumour tissue, comprising administering a chemotherapeutic agent and a
 CC composition comprising cationic liposomes consisting of cationic lipid,
 CC phosphatidylcholine and cholesterol, where oligonucleotide(s) are
 CC encapsulated within the liposome. The invention further relates to a
 CC composition comprising liposomes consisting essentially of a cationic
 CC lipid like dimethyldioctadecyl ammonium bromide (DDAB) or dimyristoyl
 CC trimethyl ammonium propane (DMTAP), phosphatidylcholine (PC),
 CC cholesterol, and containing the sequence 5'- GTGCTCCATTGATGC -3', where
 CC only the terminal sequences are phosphorothioated. The method is useful
 CC for chemosensitisation of a tumour tissue or cancer, including leukaemia,
 CC lymphoma, myeloma, carcinoma or sarcoma. The combination therapy may be
 CC used for any stage of cancer ranging from pre-cancerous lesions to cancer
 CC of advanced stages. This polynucleotide sequence represents the sense
 CC (ATG-S) raf ODN oligodeoxyribonucleotide, a cationic liposome of the
 CC invention.
 XX
 SQ Sequence 15 BP; 5 A; 4 C; 4 G; 2 T; 0 U; 0 Other;
 Query Match 100.0%; Score 15; DB 10; Length 15;
 Best Local Similarity 100.0%; Pred. No. 1.1e+02;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 1 GTGCTCCATTGATGC 15
 Db 15 GTGCTCCATTGATGC 1
 RESULT 8
 ADB97456
 ID ADB97456 standard; DNA; 15 BP.
 AC
 XX ADB97456;
 XX
 DT 04-DEC-2003 (first entry)
 XX
 DE Antisense (ATG-AS) raf ODN oligodeoxyribonucleotide.
 XX
 KM antisense; ATG-AS; raf ODN; chemosensitisation; tumour tissue;
 KM chemotherapeutic agent; cationic liposome; cationic lipid;
 KM phosphatidylcholine; cholesterol; liposome;
 KM dimethyldioctadecyl ammonium bromide; DDAB;
 KM dimyristoyl trimethyl ammonium propane; DMTAP; phosphatidylcholine; PC;
 KM cholesterol; cancer; leukaemia; lymphoma; myeloma; carcinoma; sarcoma;
 KM combination therapy; pre-cancerous lesion; chemotherapy; ss.
 XX
 OS Unidentified.
 XX
 PN WO2003070221-A1.
 XX
 PD 28-AUG-2003.
 XX
 PF 14-FEB-2003; 2003WO-US004681.
 XX
 PR 15-FEB-2002; 2002US-00075994.
 XX
 PA (GEOU) UNIV GEORGETOWN.
 PA (NEOP-) NEOPHARM INC.
 PI Kaaid U, Gokhale P, Pei J, Mewani R, Ahmad I, Drischilo A;
 PI Rahman A;
 DR WPI, 2003-689738/65.
 XX
 PT Chemosensitization of tumor tissue, useful for treating cancer, e.g.
 PT leukemia, lymphoma or myeloma, comprises administering a chemotherapeutic
 PT agent and cationic liposomes containing oligonucleotides.
 XX
 PS Example 1; Page 18; 77pp; English.

XX
 CC The invention relates to a novel method for the chemosensitisation of
 CC tumour tissue, comprising administering a chemotherapeutic agent and a
 CC composition comprising cationic liposomes consisting of cationic lipid,
 CC phosphatidylcholine and cholesterol, where oligonucleotide(s) are
 CC encapsulated within the liposome. The invention further relates to a
 CC composition comprising liposomes consisting essentially of a cationic
 CC lipid like dimethyldioctadecyl ammonium bromide (DDAB) or dimyristoyl
 CC trimethyl ammonium propane (DMTAP), phosphatidylcholine (PC),
 CC cholesterol, and containing the sequence 5'- GTGCTCCATTGATGC -3', where
 CC only the terminal sequences are phosphorothioated. The method is useful
 CC for chemosensitisation of a tumour tissue or cancer, including leukaemia,
 CC lymphoma, myeloma, carcinoma or sarcoma. The combination therapy may be
 CC used for any stage of cancer ranging from pre-cancerous lesions to cancer
 CC of advanced stages. This polynucleotide sequence represents the antisense
 CC (ATG-AS) raf ODN oligodeoxyribonucleotide, a cationic liposome of the
 CC invention.
 XX
 SQ Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;
 Query Match 100.0%; Score 15; DB 10; Length 15;
 Best Local Similarity 100.0%; Pred. No. 1.1e+02;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 1 GTGCTCCATTGATGC 15
 Db 1 GTGCTCCATTGATGC 15
 RESULT 9
 ADF82830
 ID ADF82830 standard; DNA; 15 BP.
 AC
 XX ADF82830;
 XX
 DT 26-FEB-2004 (first entry)
 XX
 DE Immunostimulant ODN25, component of lipid-nucleic acid vaccine.
 XX
 KM Immunostimulant; vaccine; lipid-nucleic acid; phosphorothioate; human;
 KM C-Raf-s; ss.
 XX
 OS Synthetic.
 OS Homo sapiens.
 XX
 FT Key Location/Qualifiers
 FT modified_base 1..15
 FT /*tag= a
 FT /mod_base= OTHER
 FT /note= "OTHER= optional phosphorothioate nucleotides"
 XX
 PN WO2003094829-A2.
 XX
 PD 20-NOV-2003.
 XX
 PF 12-MAY-2003; 2003WO-CA000680.
 XX
 PR 10-MAY-2002; 2002US-0379343P.
 PR 07-NOV-2002; 2002US-0029054S.
 PR 12-MAR-2003; 2003US-0454298P.
 XX
 PA (INEX-) INEX PHARM CORP.
 PI Semple S, Chikh G, Hope MJ, Tam YK;
 PI WPI, 2003-903935/82.
 DR WPI, 2003-903935/82.
 XX
 PT New pathogen vaccine having a lipid-nucleic acid formulation in
 PT combination with at least one microbial antigen, useful for stimulating
 PT enhanced responses against bacterial, viral and parasitic infections.
 XX
 PS Disclosure; SEQ ID NO 25; 138bp; English.
 XX

CC The present sequence is that of ODN25 (C-Raf-s) for human C-Raf-s. This
CC is an immunostimulatory oligonucleotide that can be used in lipid-nucleic
CC acid (LNA) vaccines of the invention. Claimed vaccines comprise an LNA
CC formulation in combination with at least one microbial antigen, such as
CC hepatitis B virus surface antigen. The lipid component of the LNA
CC comprises at least one cationic lipid. The oligonucleotide component of
CC the LNA preferably comprises at least one CpG dinucleotide, a methylated
CC cytosine and a phosphorothioate backbone. The vaccine is capable of
CC stimulating Th1 type humoral and cellular immune responses. An enhanced
CC humoral response is demonstrated by a strong early peak of interferon-
CC gamma production observed within hours of vaccine followed by a second
CC stronger peak of interferon-gamma production observed several days later,
CC correlated with antibody isotype switching.

CC Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;

QY Query Match 100.0%; Score 15; DB 10; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 GTGCTCCATTGATGC 15
1 GTGCTCCATTGATGC 15

RESULT 10
ADE390171

ID ADE390171 standard; DNA; 15 BP.

AC ADE390171;

DT 12-FEB-2004 (first entry)

DE Human c-raf-1 protein kinase antisense oligonucleotide.

XX ss; lipid-encapsulated therapeutic agent particle;
XX aberrant gene expression; intercellular adhesion molecule; ICAM-1; c-myc;
XX c-myc; ras; raf; erb-B-2; protein kinase C; PKC-alpha;
XX insulin-like growth factor; IGF-IR; epidermal growth factor receptor;
XX EGFR; vascular endothelial growth factor; VEGF; VEGF-R-1; tumour;
XX inflammation; infection; antisense; human.

OS Homo sapiens.

PN US2003129221-A1.

PD 10-JUL-2003.

XX 29-JUN-2001; 2001US-00895480.

XX 14-MAY-1997; 97US-00856374.

PR 14-MAY-1998; 98US-00078954.

XX (SEMP/) SEMPLE S C.
PA (KLIM/) KLIMUK S K.
PA (HARA/) HARASYM T.
PA (HOPE/) HOPE M J.
PA (ANSEL/) ANSELL S M.
PA (CULL/) CULLIS P.
PA (SCHE/) SCHERRER P.
PA (DEBE/) DEBEYER D.

PI Semple SC, Klimuk SK, Harasym T, Hope MJ, Ansell SM, Cullis P;
PI Scherrer P, Debeuyer D;
DR WPI; 2004-031296/03.

PT Preparation of a composition comprising lipid-encapsulated therapeutic
PT agent particles, useful for introducing a nucleic acid into a cell and
PT for treating diseases characterized by aberrant gene expression.

XX disclosure, SEQ ID NO 15, 52pp; English.

CC The invention relates to a method of preparation of a composition
CC comprising lipid-encapsulated therapeutic agent particles. The
CC composition is useful for introducing a nucleic acid into a cell and for
CC treating diseases characterized by aberrant gene expression (especially
CC intercellular adhesion molecule (ICAM)-1, c-myc, c-myc, ras, raf erb-B-2,
CC protein kinase C (PKC)-alpha, insulin-like growth factor (IGF)-IR,
CC epidermal growth factor receptor (EGFR), vascular endothelial growth
CC factor (VEGF) or VEGF-R-1), e.g. tumours, inflammation or infection. The
CC present sequence represents an antisense oligonucleotide.

CC Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;

QY Query Match 100.0%; Score 15; DB 12; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 GTGCTCCATTGATGC 15
1 GTGCTCCATTGATGC 15

RESULT 11
ADE39690
ID ADE39690 standard; DNA; 15 BP.

AC ADE39690;

DT 12-FEB-2004 (first entry)

DE Oligonucleotide ODN 25 (hC-Raf-1) SEQ ID NO:25.

XX cancer; vaccine; lipid-nucleic acid; LNA; tumour-associated antigen;
XX Th-1 based immune response; cytostatic; gene therapy;
XX tumour growth inhibition; tumour; human; ss.

OS Synthetic.

OS Homo sapiens.

XX Key Location/Qualifiers

FT modified_base 1..15

FT /*tag= a

FT /mod_base= OTHER

FT /note= "optionally phosphorothioate linkages"

PN WO2003094828-A2.

PD 20-NOV-2003.

XX 12-MAY-2003; 2003WO-CA000679.

XX 10-MAY-2002; 2002US-0379343P.

PR 07-NOV-2002; 2002US-00290545.

PR 04-APR-2003; 2003US-0460646P.

XX (INEX-) INEX PHARM CORP.

PA Tam YK, Semple S, Klimuk S, Chikh G;

PI WPI; 2004-011992/01.

PT New cancer vaccine having a lipid-nucleic acid formulation in combination
PT with at least one tumor-associated antigen, useful for stimulating
PT enhanced responses against tumor-associated antigens and for inhibiting
PT tumor growth.

XX Example 9, SEQ ID NO 25, 11pp; English.

CC The present invention describes a cancer vaccine (I), which comprises a
CC lipid-nucleic acid (LNA) formulation in combination with at least one
CC tumour-associated antigen that is mixed with or associated with the LNA
CC formulation comprising a lipid component having at least one cationic
CC lipid, and a nucleic acid component comprising at least one
CC oligonucleotide, where the vaccine is capable of stimulating a Th-1 based

CC immune response in vivo to the at least one tumour-associated antigen.
CC (1) has cytostatic activity, and can be used in vaccines, and in gene
CC therapy. The method and compositions of the present invention can be
CC used for stimulating enhanced responses against tumour-associated
CC antigens and for inhibiting tumour growth. The present sequence
CC represents an oligonucleotide which is used in the exemplification of the
CC present invention.
XX
SQ Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;
Query Match 100.0%; Score 15; DB 12; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GTGCTCCATTGATGC 15
Db 1 GTGCTCCATTGATGC 15

RESULT 12

ADP32025
ID ADF32025 standard; DNA; 15 BP.
XX
AC ADF32025;
XX
DT 26-FEB-2004 (first entry)
DE Antisense oligonucleotide of the invention.
XX
KM platelet; oligonucleotide; Thrombolytic; thrombocytosis; ss.
XX
OS Synthetic.
XX
PN WO2003099213-A2.
XX
PD 04-DEC-2003.
XX
PF 19-MAY-2003; 2003WO-US015922.
XX
PR 20-MAY-2002; 2002US-0382411P.
XX
PA (NEOP-) NEOPHARM INC.
XX
PI Gately ST;
XX
DR WPI; 2004-035033/03.
XX
PT Reducing the platelet count in a patient, useful for treating
PT thrombocytosis, comprises administering antisense oligonucleotides
PT inhibiting raf-1 gene with an agent that enhances penetration of the
PT oligonucleotide into cells.
XX
PS Example 1; SEQ ID NO 1; 14pp; English.
XX
CC The present invention relates to reducing the platelet count in a patient
CC comprises preparing a formulation of an oligonucleotide with an agent
CC that enhances penetration of the oligonucleotide into cells, and
CC administering the formulation to a patient having an elevated platelet
CC count. The oligonucleotide is useful for preparing a medicated platelet
CC reducing the platelet count in a patient, particularly for treating
CC thrombocytosis. The present sequence represents an oligonucleotide of the
CC invention.
XX
SQ Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;
Query Match 100.0%; Score 15; DB 12; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GTGCTCCATTGATGC 15
Db 1 GTGCTCCATTGATGC 15

RESULT 13

ADP42926
ID ADF42926 standard; DNA; 15 BP.
XX
AC ADF42926;
XX
DT 11-MAR-2004 (first entry)
DE Methylated immunostimulatory oligonucleotide ODN 25 SEQ ID NO:25.
XX
KM lipid-methylated nucleic acid formulation; immune response;
KM lipid-nucleic acid; vaccine; immunostimulant; cytostatic;
KM antiinflammatory; antiarthritic; gene therapy; cancer; inflammation;
KM arthritis; immunodeficiency disorder;
KM methylated immunostimulatory oligonucleotide; ss.
XX
OS Synthetic.
XX
PN WO2003094963-A2.
XX
PD 20-NOV-2003.
XX
PF 12-MAY-2003; 2003WO-CA000678.
XX
PR 10-MAY-2002; 2002US-0379343P.
PR 07-NOV-2002; 2002US-00290545.
PR 04-APR-2003; 2003US-0460646P.
XX
PA (INEX-) INEX PHARM CORP.
XX
PI Tam YK, Sempke S, Klimuk S, Chikh G;
XX
DR WPI; 2004-142698/14.
XX
PT lipid-methylated nucleic acid formulation for stimulating an immune
PT response in an animal comprises a lipid component and a nucleic acid
PT component comprising a methylated nucleic acid sequence.
XX
PS Disclosure; SEQ ID NO 25; 102pp; English.
XX
CC The present invention describes a lipid-methylated nucleic acid
CC formulation for stimulating an immune response in an animal, comprising a
CC lipid component and a nucleic acid component which is a methylated
CC nucleic acid sequence. Also described: (1) an adjuvant comprising a lipid
CC -nucleic acid (LNA) formulation; (2) a vaccine comprising the LNA
CC formulation in combination with at least one target antigen; (3)
CC stimulating an enhanced host immune response to antigenic stimulation,
CC comprising administering to the host the LNA formulation; (4) stimulating
CC host dendritic cells in vivo, comprising contacting at least one
CC dendritic cell with the lipid-methylated nucleic acid formulation to a
CC host; and (5) simultaneously delivering antigenic and adjuvant immune
CC stimulation to antigen presenting cells, comprising the administration of
CC the LNA formulation associated with a target antigen. The lipid-
CC methylation nucleic acid formulation has immunostimulant, cytostatic,
CC antiinflammatory and antiarthritic activities, and can be used in
CC vaccines, and in gene therapy. The formulation and methods are useful in
CC stimulating a host's immune response to antigenic stimulation, or in
CC activating and/or expanding dendritic cell populations in response to
CC antigenic stimulation. They may be used for treating cancer,
CC inflammation, arthritis or immunodeficiency disorders. The present
CC sequence represents a methylated immunostimulatory oligonucleotide given
CC in the exemplification of the present invention.
XX
SQ Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;
Query Match 100.0%; Score 15; DB 12; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GTGCTCCATTGATGC 15
Db 1 GTGCTCCATTGATGC 15

```
XX RESULT 14
XX ADL70154
XX ID ADL70154 standard; DNA; 15 BP.
XX AC ADL70154;
XX DT 20-MAY-2004 (first entry)
XX DE Oligonucleotide antisense to raf.
XX KW Raf; antisense; liposome; drug delivery; cytostatic; ss.
XX OS Synthetic.
XX PN WO2004017944-A1.
XX PD 04-MAR-2004.
XX PF 13-AUG-2003; 2003WO-US025293.
XX PR 23-AUG-2002; 2002US-0405378P.
XX PA (NEOP-) NEOPHARM INC.
XX PI Zhang J, Ahmad I;
XX WPI; 2004-257219/24.
XX PT Treatment of cellular proliferative disease e.g. cancer involves
XX PT administration of a composition comprising liposomal gemcitabine and
XX PT negatively charged phospholipid.
XX PS Disclosure; SEQ ID NO 1; 25bp; English.
XX CC The present sequence is that of an antisense oligonucleotide to raf. The
XX CC invention relates to novel gemcitabine compositions and their use in
XX CC treating proliferative diseases such as cancer, particularly in mammals,
XX CC especially in humans. The compositions include liposome-entrapped
XX CC gemcitabine. The cancer is especially lymphoma, ovarian cancer, breast
XX CC cancer, pancreatic cancer, lung cancer or colon cancer. The liposomal
XX CC gemcitabine compositions can be used in conjunction with secondary
XX CC therapeutic agents including antineoplastic, antifungal and antibiotic
XX CC agents as well as antisense oligonucleotides, especially an antisense
XX CC oligonucleotide to raf (claimed).
XX SQ Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;
XX
XX Query Match 100.0%; Score 15; DB 12; Length 15;
XX Best Local Similarity 100.0%; Pred. No. 1.1e+02;
XX Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 GTGCTCCATTGATGC 15
XX | | | | | | | | | |
XX Db 1 GTGCTCCATTGATGC 15
XX
XX RESULT 15
XX ADR88950
XX ID ADR88950 standard; DNA; 15 BP.
XX AC ADR88950;
XX DT 18-NOV-2004 (first entry)
XX DE Anti c-raf-1 oligonucleotide.
XX KW C-raf-1; liposomal; antineoplastic; cytostatic; cancer; antisense; ss.
XX OS Synthetic.
XX PN WO2004071466-A2.
```

```
XX PD 26-AUG-2004.
XX PF 11-FEB-2004; 2004WO-US004555.
XX PR 11-FEB-2003; 2003US-0446895P.
XX PA (NEOP-) NEOPHARM INC.
XX PI Bhamidipati S, Ahmad Z, Ahmad I;
XX WPI; 2004-635030/61.
XX PT Preparation of liposomal composition used for treating e.g. cancer
XX PT involves dissolving lipid fraction in water miscible organic solvent and
XX PT mixing solvent solution with aqueous solution.
XX PS Disclosure; Page 6; 27pp; English.
XX CC The invention relates to the preparation of a liposomal composition. The
XX CC method involves: dissolving a lipid fraction in a water-miscible organic
XX CC solvent; and mixing the water-miscible organic solvent solution
XX CC comprising the lipid fraction with an aqueous solution under conditions
XX CC to form a bulk liposomal composition. The method further involves adding
XX CC at least one active principal to the water-miscible organic solvent prior
XX CC to the addition of the lipid fraction, or to the aqueous solution prior,
XX CC during or after the step (b), size-reducing the bulk liposomal
XX CC composition to obtain a size-reduced liposomal composition, freeing the
XX CC liposomal composition of the water-miscible organic solvent by
XX CC diafiltration using a tangential flow filtration process and sterile
XX CC filtration, sterile-filtering the liposomal composition and freeze-drying
XX CC the liposomal preparation. Step (b) involves adding the water-miscible
XX CC organic solvent solution to the aqueous solution while mixing and mixing
XX CC of solution following addition of water-miscible solvent comprising the
XX CC lipid fraction to the aqueous solution while cooling. The active
XX CC principal comprises at least one antineoplastic or antifungal agent
XX CC (preferably taxane, camptothecin or their derivatives, especially
XX CC paclitaxel or docetaxel). The composition is used for the treatment of
XX CC disease e.g. cancer. The composition eliminates the disease or its
XX CC symptoms, need not completely eradicate the effects of the disease,
XX CC reduces the severity of a disease, infection or reduction in the rate by
XX CC which a disease progresses within a patient. The method permits the
XX CC production of liposomal formulation on a commercial scale. The present
XX CC sequence represents an antisense oligonucleotide specific for c-raf-1,
XX CC that can be used as an active principal.
XX SQ Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;
XX
XX Query Match 100.0%; Score 15; DB 13; Length 15;
XX Best Local Similarity 100.0%; Pred. No. 1.1e+02;
XX Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 GTGCTCCATTGATGC 15
XX | | | | | | | | | |
XX Db 1 GTGCTCCATTGATGC 15
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OM nucleic - nucleic search, using sw model

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Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

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1: gb_ha:*
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14: gb_vl:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	15	100.0	15	6	AR110775 Sequence
2	15	100.0	15	6	AR110777 Sequence
3	15	100.0	15	6	AR167449 Sequence
4	15	100.0	15	6	CQ789712 Sequence
5	15	100.0	15	6	AR310685 Sequence
6	15	100.0	15	6	AR310687 Sequence
7	15	100.0	15	6	AX979662 Sequence
8	15	100.0	15	6	AX957646 Sequence
9	15	100.0	15	6	AX957740 Sequence
10	15	100.0	15	6	AX958145 Sequence
11	15	100.0	15	6	BD106498 Sequence
12	15	100.0	20	6	AR073978 Sequence
13	15	100.0	20	6	AR216002 Sequence
14	15	100.0	25	6	AR110776 Sequence
15	15	100.0	25	6	AR310686 Sequence
16	14	93.3	35	10	AB112659S1
17	13	86.7	20	6	AR073933 Sequence
18	13	86.7	20	6	AR105501 Sequence
19	13	86.7	20	6	E49512 Antisense O

20	13	86.7	20	6	127232
21	13	86.7	20	6	AR215955
22	12.4	82.7	20	6	AR037100
23	12.4	82.7	20	6	AR070338
24	12.4	82.7	20	6	AX294189
25	12.4	82.7	21	6	AX598466
26	12.4	82.7	24	6	AX289556
27	12.4	82.7	27	6	AR039324
28	12.4	82.7	43	6	C0749113
29	12	80.0	20	6	AR073934
30	12	80.0	20	6	AR106990
31	12	80.0	20	6	AR106991
32	12	80.0	20	6	E49513
33	12	80.0	20	6	127233
34	12	80.0	20	6	AR215956
35	12	80.0	22	6	AR493387
36	11.8	78.7	26	6	A16281
37	11.8	78.7	27	6	A16267
38	11.8	78.7	27	6	A16267
39	11.8	78.7	27	6	AR080410
40	11.8	78.7	27	6	AR092534
41	11.8	78.7	27	6	AR122889
42	11.8	78.7	27	6	AR123544
43	11.8	78.7	27	6	AR148361
44	11.8	78.7	30	6	AR069912
45	11.8	78.7	33	6	BD014421

ALIGNMENTS

RESULT 1	AR110775	Sequence 1 from patent US 6126965.	15 bp	DNA	linear	PAT 14-FEB-2001
LOCUS	AR110775					
DEFINITION	Sequence 1 from patent US 6126965.					
ACCESSION	AR110775					
VERSION	AR110775.1	GI:12827623				
KEYWORDS						
SOURCE	Unknown.					
ORGANISM	Unknown.					
REFERENCE	1 (bases 1 to 15)					
AUTHORS	Kasid,U., Gokhale,P., Dritschilo,A. and Rahman,A.					
TITLE	Liposomes containing oligonucleotides					
JOURNAL	Patent: US 6126965-A 1 03-Oct-2000;					
FEATURES	Location/Qualifiers					
source	1..15					
ORIGIN	/organism="unknown"					
	/mol_type="unassigned DNA"					
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Best Local Similarity	100.0%; Pred. No. 1.2e+03;					
Matches	15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;					
Qy	1 GTGCTCCATTGATGC 15					
Db	1 GTGCTCCATTGATGC 15					
RESULT 2	AR110777/c	Sequence 3 from patent US 6126965.	15 bp	DNA	linear	PAT 14-FEB-2001
LOCUS	AR110777					
DEFINITION	Sequence 3 from patent US 6126965.					
ACCESSION	AR110777					
VERSION	AR110777.1	GI:12827625				
KEYWORDS						
SOURCE	Unknown.					
ORGANISM	Unknown.					
REFERENCE	1 (bases 1 to 15)					
AUTHORS	Kasid,U., Gokhale,P., Dritschilo,A. and Rahman,A.					
TITLE	Liposomes containing oligonucleotides					

JOURNAL Patent: US 6126965-A 3 03-OCT-2000;
FEATURES Location/Qualifiers
source 1..15
/organism="unknown"
/mol_type="unassigned DNA"

ORIGIN

Query Match 100.0%; Score 15; DB 6; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.2e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTGCTCCATTGATGC 15
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15 GTGCTCCATTGATGC 1

Db 15 GTGCTCCATTGATGC 1

RESULT 3
LOCUS AR167449 15 bp DNA linear PAT 17-DEC-2001
DEFINITION Sequence 15 from patent US 6287591.
ACCESSION AR167449
VERSION AR167449.1 GI:17903229
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 15)
AUTHORS Sample,S.C., Klimuk,S.K., Harasym,T., Hope,M.J., Ansell,S.M.,
Challis,P., Scherrer,P. and Debeyer,D.,
Charged therapeutic agents encapsulated in lipid particles
containing four lipid components
Patent: US 6287591-A 15 11-SEP-2001;
location/Qualifiers
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/organism="unknown"
/mol_type="unassigned DNA"

ORIGIN

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Best Local Similarity 100.0%; Pred. No. 1.2e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTGCTCCATTGATGC 15
|||||
1 GTGCTCCATTGATGC 15

Db 1 GTGCTCCATTGATGC 15

RESULT 4
LOCUS CQ789712 15 bp DNA linear PAT 29-MAR-2004
DEFINITION Sequence 1 from Patent WO2004017944.
ACCESSION CQ789712
VERSION CQ789712.1 GI:45823264
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
REFERENCE 1
AUTHORS Zhang,J.A. and Ahmad,I.
TITLE Liposomal gemcitabine compositions for better drug delivery
JOURNAL Patent: WO 2004017944-A 1 04-MAR-2004;
Neopharm, Inc. (US)
location/Qualifiers
1..15
/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="Anti-rai-oligonucleotides"

ORIGIN

Query Match 100.0%; Score 15; DB 6; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.2e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTGCTCCATTGATGC 15
|||||
1 GTGCTCCATTGATGC 15

Db 1 GTGCTCCATTGATGC 15

RESULT 5
LOCUS AR310685 15 bp DNA linear PAT 12-JUN-2003
DEFINITION Sequence 1 from patent US 6559129.
ACCESSION AR310685
VERSION AR310685.1 GI:31703829
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 15)
AUTHORS Kasid,U., Gokhale,P., Zhang,C., Dritschilo,A. and Rahman,A.
TITLE Cationic liposomal delivery system and therapeutic use thereof
JOURNAL Patent: US 6559129-A 1 06-MAY-2003;
location/Qualifiers
1..15
/organism="unknown"
/mol_type="genomic DNA"

ORIGIN

Query Match 100.0%; Score 15; DB 6; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.2e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTGCTCCATTGATGC 15
|||||
1 GTGCTCCATTGATGC 15

Db 1 GTGCTCCATTGATGC 15

RESULT 6
LOCUS AR310687/c 15 bp DNA linear PAT 12-JUN-2003
DEFINITION Sequence 3 from patent US 6559129.
ACCESSION AR310687
VERSION AR310687.1 GI:31703831
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 15)
AUTHORS Kasid,U., Gokhale,P., Zhang,C., Dritschilo,A. and Rahman,A.
TITLE Cationic liposomal delivery system and therapeutic use thereof
JOURNAL Patent: US 6559129-A 3 06-MAY-2003;
location/Qualifiers
1..15
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/mol_type="genomic DNA"

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Query Match 100.0%; Score 15; DB 6; Length 15;
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Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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15 GTGCTCCATTGATGC 1

Db 15 GTGCTCCATTGATGC 1

RESULT 7
LOCUS AX797662 15 bp DNA linear PAT 04-OCT-2003
DEFINITION Sequence 25 from Patent WO03039595.
ACCESSION AX797662
VERSION AX797662.1 GI:37518090
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
TITLE 1
JOURNAL Sample, S., Klimuk, S. and Yuan, Z.N.
Mucosal adjuvants comprising an oligonucleotide and a cationic
lipid
Patent: WO 03039595-A 25 15-MAY-2003;
Inex Pharmaceuticals Corp. (CA)
FEATURES location/Qualifiers
SOURCE 1.15
/organism="Homo sapiens"
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/db_xref="taxon:9606"
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Best Local Similarity 100.0%; Pred. No. 1.2e+03;
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DB 1 GTGCTCCATTGATGC 15

RESULT 8
LOCUS AX957646 15 bp DNA linear PAT 08-JAN-2004
DEFINITION Sequence 25 from Patent WO03094963.
ACCESSION AX957646
VERSION AX957646.1 GI:40785518
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE 1
AUTHORS Tan, Y.K., Semple, S., Klimuk, S. and Chikh, G.
TITLE Methylated immunostimulatory oligonucleotides and methods of using
JOURNAL the same
Patent: WO 03094963-A 25 20-NOV-2003;
Inex Pharmaceuticals Corporation (CA)
FEATURES location/Qualifiers
SOURCE 1.15
/organism="Homo sapiens"
/mol_type="unassigned DNA"
/db_xref="taxon:9606"
ORIGIN

Query Match 100.0%; Score 15; DB 6; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.2e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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DB 1 GTGCTCCATTGATGC 15

RESULT 9
LOCUS AX957740 15 bp DNA linear PAT 08-JAN-2004
DEFINITION Sequence 25 from Patent WO03094828.
ACCESSION AX957740
VERSION AX957740.1 GI:40785558
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE 1
AUTHORS Tam, Y.K., Semple, S., Klimuk, S. and Chikh, G.
TITLE Cancer vaccines and methods of using the same
JOURNAL Patent: WO 03094828-A 25 20-NOV-2003;
Inex Pharmaceuticals Corp. (CA)

FEATURES location/Qualifiers
SOURCE 1.15
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/mol_type="unassigned DNA"
/db_xref="taxon:9606"
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DB 1 GTGCTCCATTGATGC 15

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LOCUS AX958145 15 bp DNA linear PAT 08-JAN-2004
DEFINITION Sequence 25 from Patent WO03094829.
ACCESSION AX958145
VERSION AX958145.1 GI:40785809
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE 1
AUTHORS Semple, S., Chikh, G., Hope, M.J. and Tam, Y.K.
TITLE Pathogen vaccines and methods for using the same
JOURNAL Patent: WO 03094829-A 25 20-NOV-2003;
Inex Pharmaceuticals Corp. (CA)
FEATURES location/Qualifiers
SOURCE 1.15
/organism="Homo sapiens"
/mol_type="unassigned DNA"
/db_xref="taxon:9606"
ORIGIN

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Best Local Similarity 100.0%; Pred. No. 1.2e+03;
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OY 1 GTGCTCCATTGATGC 15
DB 1 GTGCTCCATTGATGC 15

RESULT 11
LOCUS BD106498 15 bp DNA linear PAT 18-SEP-2002
DEFINITION High efficiency encapsulation of charged therapeutic agents in
lipid vesicles.
ACCESSION BD106498
VERSION BD106498.1 GI:23201316
KEYWORDS JP 2002501511-A/15.
SOURCE Chlamydia sp.
ORGANISM Chlamydia
REFERENCE 1 (bases 1 to 15)
AUTHORS Semple, S.C., Klimuk, S.K., Harasym, T., Hope, M.J., Ansel, S.M.,
Cullis, P., Scherrer, P. and Debever, D.S.
TITLE High efficiency encapsulation of charged therapeutic agents in
lipid vesicles
JOURNAL Patent: JP 2002501511-A 15 15-JAN-2002;
INEX PHARMACEUTICALS CORP
COMMENT JP 2002501511-A/15
PD 15-JAN-2002
PF 14-MAY-1998 JP 1998548646
PI SEAN C SEMPLE, SANDRA K KLIMUK, TROY HARASYM, MICHAEL J HOPE, PI
PIETER CULLIS, PETER SCHERRER, DAN SUITE DEBEVER PC A61K9/00
CC Strandedness: Single;

CC Topology: linear;
FH Key Location/Qualifiers.
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/db_xref="taxon:35827"

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1 GTGCTCCATTGATGC 15

RESULT 12
AR073978 20 bp DNA linear PAT 28-AUG-2000
LOCUS AR073978
DEFINITION Sequence 47 from patent US 5952229.
ACCESSION AR073978
VERSION AR073978.1 GI:10000738
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 20)
AUTHORS Monia,B.P. and Boggs,R.T.
TITLE Antisense oligonucleotide modulation of raf gene expression
JOURNAL Patent: US 5952229-A 47 14-SEP-1999;
FEATURES
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/mol_type="unassigned DNA"

ORIGIN
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Best Local Similarity 100.0%; Pred. No. 1.1e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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5 GTGCTCCATTGATGC 19

RESULT 13
AR216002 20 bp DNA linear PAT 25-SEP-2002
LOCUS AR216002
DEFINITION Sequence 49 from patent US 6410518.
ACCESSION AR216002
VERSION AR216002.1 GI:23314290
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 20)
AUTHORS Monia,B.P.
TITLE Antisense oligonucleotide inhibition of raf gene expression
JOURNAL Patent: US 6410518-A 49 25-JUN-2002;
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/mol_type="genomic DNA"

ORIGIN
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Best Local Similarity 100.0%; Pred. No. 1.1e+03;
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1 GTGCTCCATTGATGC 15

DB 5 GTGCTCCATTGATGC 19

RESULT 14
AR110776 25 bp DNA linear PAT 14-FEB-2001
LOCUS AR110776
DEFINITION Sequence 2 from patent US 6126965.
ACCESSION AR110776
VERSION AR110776.1 GI:12827624
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 25)
AUTHORS Kasid,U., Gokhale,P., Dritschilo,A. and Rahman,A.
TITLE Liposomes containing oligonucleotides
JOURNAL Patent: US 6126965-A 2 03-OCT-2000;
FEATURES
source 1..25
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ORIGIN
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Best Local Similarity 100.0%; Pred. No. 1.1e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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RESULT 15
AR310686 25 bp DNA linear PAT 12-JUN-2003
LOCUS AR310686
DEFINITION Sequence 2 from patent US 6559129.
ACCESSION AR310686
VERSION AR310686.1 GI:31703830
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 25)
AUTHORS Kasid,U., Gokhale,P., Zhang,C., Dritschilo,A. and Rahman,A.
TITLE Cationic liposomal delivery system and therapeutic use thereof
JOURNAL Patent: US 6559129-A 2 06-MAY-2003;
FEATURES
source 1..25
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ORIGIN
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Best Local Similarity 100.0%; Pred. No. 1.1e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTGCTCCATTGATGC 15
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Search completed: June 21, 2005, 06:00:54
Job time : 1689 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: June 21, 2005, 06:01:03 ; Search time 6157 Seconds
(without alignments)
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Title: US-10-075-994A-1

Perfect score: 15
Sequence: 1 gtcgtcatcgtatgc 15

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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2	15	100.0	15	10	US-09-930-283A-3
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4	15	100.0	15	15	US-10-290-545-25
5	15	100.0	15	16	US-10-365-623-16
6	15	100.0	15	17	US-10-075-994A-1
7	15	100.0	15	17	US-10-075-994A-3

8	15	100.0	15	17	US-10-075-994A-4	Sequence 4, Appli
9	15	100.0	15	17	US-10-347-924-1	Sequence 1, Appli
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24	13	86.7	25	21	US-10-719-900-495729	Sequence 495729,
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26	12.4	82.7	21	17	US-10-321-039-657	Sequence 657, App
27	12.4	82.7	25	21	US-10-719-900-12019	Sequence 12019, A
28	12.4	82.7	25	21	US-10-719-900-59921	Sequence 59921, A
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34	12.4	82.7	25	21	US-10-719-900-781945	Sequence 781945,
35	12.4	82.7	25	21	US-10-809-189-36748	Sequence 36748, A
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43	12	80.0	20	14	US-10-280-600-6	Sequence 6, Appli
44	12	80.0	20	14	US-10-281-312-5	Sequence 5, Appli
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ALIGNMENTS

RESULT 1
US-09-930-283A-1
; Sequence 1, Application US/09930283A
; Patent No. US20020160038A1
GENERAL INFORMATION:
APPLICANT: Kasid, Usha
Gokhale, Prafulla
Ditechilo, Anatoly
Rahman, Agulur
TITLE OF INVENTION: Liposomes containing Oligonucleotides
NUMBER OF SEQUENCES: 3
CORRESPONDENCE ADDRESS:
ADDRESSEE: Hendricks and Assoc.
STREET: P.O. Box 2509
CITY: Fairfax
STATE: VA
COUNTRY: US
ZIP: 22031
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/930, 283A
FILING DATE: 16-Aug-2001
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 09/354, 109
FILING DATE: 1999-07-15

ATTORNEY/AGENT INFORMATION:
NAME: Hendricks, Glenna
REGISTRATION NUMBER: 32,535
TELECOMMUNICATION INFORMATION:
TELEPHONE: (703) 591-4470
TELEFAX: (703) 591-4428
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 15 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: unknown
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: YES
SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-09-930-283A-1

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Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 1 GTGCTCCATTGATGC 15

RESULT 2
US-09-930-283A-3/C
Sequence 3, Application US/09930283A
Patent No. US20020160038A1
GENERAL INFORMATION:
APPLICANT: Kasid, Usha
Gokhale, Prafulla
Dritschilo, Anatoly
Rahman, Aquilur
TITLE OF INVENTION: Liposomes containing Oligonucleotides
NUMBER OF SEQUENCES: 3
CORRESPONDENCE ADDRESS:
ADDRESSEE: Hendricks and Assoc.
STREET: P.O. Box 2509
CITY: Fairfax
STATE: VA
COUNTRY: US
ZIP: 22031
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/930,283A
FILING DATE: 16-Aug-2001
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 09/354,109
FILING DATE: 1999-07-15
ATTORNEY/AGENT INFORMATION:
NAME: Hendricks, Glenna
REGISTRATION NUMBER: 32,535
REFERENCE/DOCKET NUMBER: Kasid
TELECOMMUNICATION INFORMATION:
TELEPHONE: (703) 591-4470
TELEFAX: (703) 591-4428
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 15 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: unknown
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO

ANTI-SENSE: YES
SEQUENCE DESCRIPTION: SEQ ID NO: 3:
US-09-930-283A-3

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Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 15 GTGCTCCATTGATGC 1

RESULT 3
US-09-895-480A-15
Sequence 15, Application US/09895480A
Publication No. US20030129221A1
GENERAL INFORMATION:
APPLICANT: Inex Pharmaceuticals Inc.
TITLE OF INVENTION: High Efficiency Encapsulation of Charged Therapeutic Agents in Lipid Vesicles
NUMBER OF SEQUENCES: 17
CORRESPONDENCE ADDRESS:
ADDRESSEE: Oppedahl & Larson LLP
STREET: PO Box 5068
CITY: Dillon
STATE: CO
COUNTRY: US
ZIP: 80435
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.5 inch, 1.44 MB
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS 5.0
SOFTWARE: Word Perfect
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/895,480A
FILING DATE: 29-Jun-2001
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: <Unknown>
FILING DATE: <Unknown>
ATTORNEY/AGENT INFORMATION:
NAME: <Unknown>
REGISTRATION NUMBER: <Unknown>
REFERENCE/DOCKET NUMBER: <Unknown>
TELECOMMUNICATION INFORMATION:
TELEPHONE: <Unknown>
TELEFAX: <Unknown>
INFORMATION FOR SEQ ID NO: 15:
SEQUENCE CHARACTERISTICS:
LENGTH: 15
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
HYPOTHETICAL: no
ANTI-SENSE: yes
SEQUENCE DESCRIPTION: SEQ ID NO: 15:
US-09-895-480A-15

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Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 1 GTGCTCCATTGATGC 15

RESULT 4
US-10-290-545-25

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: Sequence 25 Application US/10290545
: Publication No. US20030125292A1
: GENERAL INFORMATION:
: APPLICANT: Semple, Sean
: APPLICANT: Klimuk, Sandy
: APPLICANT: Yuan, Zuan-Ning
: TITLE OF INVENTION: Improved Mucosal Vaccines and Methods for Using the Same
: FILE REFERENCE: A-71854/PA1/AXG
: CURRENT APPLICATION NUMBER: US/10/290,545
: CURRENT FILING DATE: 2002-11-07
: NUMBER OF SEQ ID NOS: 30
: SOFTWARE: PatentIn version 3.1
: SEQ ID NO 25
: LENGTH: 15
: TYPE: DNA
: ORGANISM: Homo sapiens
US-10-290-545-25

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Query Match	100.0%;	Score 15;	DB 15;	Length 15;
Best Local Similarity	100.0%;	Pred. No. 1.3e+02;		
Matches 15;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;

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Db	1	GTGCTCCATTGATGC	15

RESULT 5

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? Sequence 16 Application US/10365623
? Publication No. US20030166512A1
? GENERAL INFORMATION:
? APPLICANT: Xie, Dong
? TITLE OF INVENTION: Protein Carrier System for Therapeutic Oligonucleotides
? FILE REFERENCE: 63024.000001
? CURRENT APPLICATION NUMBER: US/10/365,623
? CURRENT FILING DATE: 2003-02-13
? NUMBER OF SEQ ID NOS: 23
? SOFTWARE: PatentIn version 3.1
? SEQ ID NO 16
? LENGTH: 15
? TYPE: DNA
? ORGANISM: Artificial Sequence
? FEATURE:
? OTHER INFORMATION: Antisense oligonucleotide directed against human c-rac-1 protein
? OS-10-365-623-16

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Query Match	100.0%	Score 15;	DB 16;	Length 15;
Best Local Similarity	100.0%	Pred. No. 1.3e+02;		
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RESULT 6

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US/10-075-994A-1
: Sequence 1, Application US/10075994A
: Publication No. US20030215489A1
: GENERAL INFORMATION:
: APPLICANT: KASID, Usha
: APPLICANT: GOKHALE, Prafulla
: APPLICANT: PIE, Jin
: APPLICANT: MEWANI, Rajshree
: APPLICANT: AHMAD, Imran
: APPLICANT: DRITSCHILLO, Anatoliy
: APPLICANT: RAHMAN, Aquilur
: TITLE OF INVENTION: CHEMOSENSITIZING WITH LIPOSOMES CONTAINING OLIGONUCLEOTIDES
: FILE REFERENCE: 219604
: CURRENT APPLICATION NUMBER: US/10/075,994A
: CURRENT FILING DATE: 2002-02-15

```

US-10-075-994A-1

Query Match	100.0%;	Score 15;	DB 17;	Length 15;
Best Local Similarity	100.0%;	Pred. No. 1.3e+02;		
Matches	15;	Conservative	0;	Mismatches 0;
				Indels 0;
				Gaps 0;

```
QY      1 GTGCTCCATTGATGC 15
         |||||
Db      1 GTGCTCCATTGATGC 15
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RESULT 7
US-10-075-994A-3/c

```

Sequence 3, Application US/10/075994A
Publication No. US20030215489A1
GENERAL INFORMATION:
APPLICANT: KASID, Usha
APPLICANT: GOKHALE, Prafulla
APPLICANT: PTE, Jin
APPLICANT: MEMANI, Rajshree
APPLICANT: AHMAD, Imran
APPLICANT: DRITSCHLO, Anatoly
APPLICANT: RAHMANN, Aquilur
TITLE OF INVENTION: CHEMOSENSITIZING WITH LIPOSOMES CONTAINING OLIGONUCLEOTIDES
FILE REFERENCE: 219604
CURRENT APPLICATION NUMBER: US/10/075,994A
CURRENT FILING DATE: 2002-02-15
NUMBER OF SEQ ID NOS: 7
SOFTWARE: PatentIn version 3.2
SEQ ID NO 3
LENGTH: 15
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Synthetic
US-10-075-994A-3

```

Query Match	100.0%	Score 15;	DB 17;	Length 15;
Best Local Similarity	100.0%	Pred. No. 1.3e+02;		
Matches 15;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;

```

QY      1 GTGCTCCATTGATGC 15
          |||||
Db      15 GTGCTCCATTGATGC 1

```

RESULT 8

```

/ Sequence 4, Application US/10075994A
/ US-10-075-994A-4
/ Publication No. US20030215489A1
/ GENERAL INFORMATION:
/ APPLICANT: KASID, Usha
/ APPLICANT: GOKHALE, Prafulla
/ APPLICANT: PIE, Jin
/ APPLICANT: MEWANI, Rajshree
/ APPLICANT: AHMAD, Imran
/ APPLICANT: DRITSCHILLO, Anatoly
/ APPLICANT: RAHMAN, Aguilur
/ TITLE OF INVENTION: CHEMOSENSITIZING WITH LIPOSOMES CONTAINING OLIGONUCLEOTIDES
/ FILE REFERENCE: 219604
/ CURRENT APPLICATION NUMBER: US/10/075,994A
/ CURRENT FILING DATE: 2002-02-15
/ NUMBER OF SEQ ID NOS: 7
/ SOFTWARE: PatentIn version 3.2
/ SEQ ID NO 4

```

LENGTH: 15
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Synthetic
US-10-075-994a-4

Query Match 100.0%; Score 15; DB 17; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GTGCTCCATTGATGC 15
Db 1 GTGCTCCATTGATGC 15

RESULT 9
US-10-347-924-1
Sequence 1, Application US/10347924
Publication No. US20030229040A1
GENERAL INFORMATION:
APPLICANT: Kasid, Usha
APPLICANT: Gokhale, Prafulla
APPLICANT: Zhang, Chuando
APPLICANT: Dristschillo, Anatoly
APPLICANT: Rahman, Aquilur
TITLE OF INVENTION: CATIONIC LIPOSOMAL DELIVERY SYSTEM AND THERAPEUTIC USE THEREOF
FILE REFERENCE: 220807
CURRENT APPLICATION NUMBER: US/10/347,924
CURRENT FILING DATE: 2003-01-21
PRIOR APPLICATION NUMBER: US 09/354,109
PRIOR FILING DATE: 1999-07-15
PRIOR APPLICATION NUMBER: US 08/957,327
PRIOR FILING DATE: 1997-10-24
PRIOR APPLICATION NUMBER: US 60/041,192
PRIOR FILING DATE: 1997-03-21
NUMBER OF SEQ ID NOS: 1
SOFTWARE: PatentIn version 3.2
SEQ ID NO 1
LENGTH: 15
TYPE: DNA
ORGANISM: Artificial
FEATURE:
OTHER INFORMATION: Oligonucleotide
US-10-347-924-1

Query Match 100.0%; Score 15; DB 17; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GTGCTCCATTGATGC 15
Db 1 GTGCTCCATTGATGC 15

RESULT 10
US-10-437-263-25
Sequence 25, Application US/10437263
Publication No. US20040009943A1
GENERAL INFORMATION:
APPLICANT: Semple, Sean
APPLICANT: Tam, Ying K.
APPLICANT: Chikh, Ghania
APPLICANT: Hope, Michael J.
TITLE OF INVENTION: PATHOGEN VACCINES AND METHODS FOR USING THE SAME
FILE REFERENCE: A-72216/TAL
CURRENT APPLICATION NUMBER: US/10/437,263
CURRENT FILING DATE: 2003-05-12
PRIOR APPLICATION NUMBER: 60/379,343
PRIOR FILING DATE: 2002-05-10
PRIOR APPLICATION NUMBER: 60/460,646
PRIOR FILING DATE: 2003-04-04
PRIOR APPLICATION NUMBER: 60/454,298

PRIOR FILING DATE: 2003-03-12
NUMBER OF SEQ ID NOS: 34
SOFTWARE: PatentIn version 3.2
SEQ ID NO 25
LENGTH: 15
TYPE: DNA
ORGANISM: Homo sapiens
US-10-437-263-25

Query Match 100.0%; Score 15; DB 17; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GTGCTCCATTGATGC 15
Db 1 GTGCTCCATTGATGC 15

RESULT 11
US-10-437-275-25
Sequence 25, Application US/10437275
Publication No. US20040009944A1
GENERAL INFORMATION:
APPLICANT: Tam, Ying K.
APPLICANT: Semple, Sean
APPLICANT: Klimuk, Sandra
APPLICANT: Chikh, Ghania
TITLE OF INVENTION: METHYLATED IMMUNOSTIMULATORY OLIGONUCLEOTIDES AND METHODS OF
FILE REFERENCE: A-72158/TAL
CURRENT APPLICATION NUMBER: US/10/437,275
CURRENT FILING DATE: 2003-05-12
PRIOR APPLICATION NUMBER: 60/379,343
PRIOR FILING DATE: 2002-05-10
PRIOR APPLICATION NUMBER: 60/460,646
PRIOR FILING DATE: 2003-04-04
PRIOR APPLICATION NUMBER: 10/290,545
PRIOR FILING DATE: 2002-11-07
NUMBER OF SEQ ID NOS: 32
SOFTWARE: PatentIn version 3.2
SEQ ID NO 25
LENGTH: 15
TYPE: DNA
ORGANISM: Homo sapiens
US-10-437-275-25

Query Match 100.0%; Score 15; DB 17; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GTGCTCCATTGATGC 15
Db 1 GTGCTCCATTGATGC 15

RESULT 12
US-10-437-258-25
Sequence 25, Application US/10437258
Publication No. US20040013649A1
GENERAL INFORMATION:
APPLICANT: Tam, Ying K.
APPLICANT: Semple, Sean
APPLICANT: Klimuk, Sandra
APPLICANT: Chikh, Ghania
TITLE OF INVENTION: CANCER VACCINES AND METHODS OF USING THE SAME
FILE REFERENCE: A-72252/TAL
CURRENT APPLICATION NUMBER: US/10/437,258
CURRENT FILING DATE: 2003-05-12
PRIOR APPLICATION NUMBER: 60/379,343
PRIOR FILING DATE: 2002-05-10
PRIOR APPLICATION NUMBER: 60/460,646
PRIOR FILING DATE: 2003-04-04
PRIOR APPLICATION NUMBER: 60/454,298

PRIOR FILING DATE: 2003-03-12
NUMBER OF SEQ ID NOS: 34
SOFTWARE: PatentIn version 3.2
SEQ ID NO 25
LENGTH: 15
TYPE: DNA
ORGANISM: Homo sapiens
US-10-437-258-25

Query Match 100.0%; Score 15; DB 17; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GTGCTCATTTGATGC 15
Db 1 GTGCTCATTTGATGC 15

RESULT 13

US-10-925-734-15
Sequence 15, Application US/10925734
Publication No. US2005008689A1
GENERAL INFORMATION:

APPLICANT: Inex Pharmaceuticals Inc.
TITLE OF INVENTION: High Efficiency Encapsulation of Charged
Therapeutic Agents in Lipid Vesicles

NUMBER OF SEQUENCES: 17

CORRESPONDENCE ADDRESS:

ADDRESSEE: Oppedahl & Larson LLP

STREET: PO Box 5068

CITY: Dillon

STATE: CO

COUNTRY: US

ZIP: 80435

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette, 3.5 inch, 1.44 Mb

COMPUTER: IBM Compatible

OPERATING SYSTEM: DOS 5.0

SOFTWARE: Word Perfect

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/10/925,734

FILING DATE: 24-Aug-2004

CLASSIFICATION: <Unknown>

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US/09/895,480

FILING DATE: 29-Jun-2001

ATTORNEY/AGENT INFORMATION:

NAME: <Unknown>

REGISTRATION NUMBER: <Unknown>

REFERENCE/DOCKET NUMBER: <Unknown>

TELECOMMUNICATION INFORMATION:

TELEPHONE: <Unknown>

TELEFAX: <Unknown>

TELEX: <Unknown>

INFORMATION FOR SEQ ID NO: 15:

SEQUENCE CHARACTERISTICS:

LENGTH: 15

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: other nucleic acid

HYPOTHETICAL: no

ANTI-SENSE: yes

SEQUENCE DESCRIPTION: SEQ ID NO: 15:

US-10-925-734-15

OY 1 GTGCTCATTTGATGC 15

Db 1 GTGCTCATTTGATGC 15

RESULT 14

US-10-057-550-49
Sequence 49, Application US/10057550
Publication No. US20030032607A1
GENERAL INFORMATION:

APPLICANT: Monia, Brett P.
TITLE OF INVENTION: Antisense Oligonucleotide Modulation of raf Gene Expression
FILE REFERENCE:

CURRENT APPLICATION NUMBER: US/10/057,550

CURRENT FILING DATE: 2002-01-25

PRIOR APPLICATION NUMBER: 09/506,073

PRIOR FILING DATE: 2000-02-18

PRIOR APPLICATION NUMBER: US 09/143,214

PRIOR FILING DATE: 1998-08-28

PRIOR APPLICATION NUMBER: PCT/US98/13961

PRIOR FILING DATE: 1998-07-06

PRIOR APPLICATION NUMBER: US 08/888,982

PRIOR FILING DATE: 1997-07-07

PRIOR APPLICATION NUMBER: US 08/756,806

PRIOR FILING DATE: 1996-11-26

PRIOR APPLICATION NUMBER: PCT/US95/07111

PRIOR FILING DATE: 1995-05-31

PRIOR APPLICATION NUMBER: US 08/250,856

PRIOR FILING DATE: 1994-05-31

NUMBER OF SEQ ID NOS: 130

SEQ ID NO 49

LENGTH: 20

TYPE: DNA

ORGANISM: artificial sequence

FEATURE:

OTHER INFORMATION: antisense sequence

US-10-057-550-49

OY 1 GTGCTCATTTGATGC 15

Db 5 GTGCTCATTTGATGC 19

RESULT 15

US-10-173-225B-47
Sequence 47, Application US/10173225B
Publication No. US20030119769A1
GENERAL INFORMATION:

APPLICANT: Monia, Brett P.
TITLE OF INVENTION: Antisense Oligonucleotide Modulation of raf Gene Expression
FILE REFERENCE: ISPH-0665

CURRENT APPLICATION NUMBER: US/10/173,225B

CURRENT FILING DATE: 2002-12-06

PRIOR APPLICATION NUMBER: US 10/057,550

PRIOR FILING DATE: 2002-01-25

PRIOR APPLICATION NUMBER: US 09/143,214

PRIOR FILING DATE: 1998-08-28

PRIOR APPLICATION NUMBER: PCT/US98/13961

PRIOR FILING DATE: 1998-07-06

PRIOR APPLICATION NUMBER: US 08/888,982

PRIOR FILING DATE: 1997-07-07

PRIOR APPLICATION NUMBER: US 08/756,806

PRIOR FILING DATE: 1996-11-26

PRIOR APPLICATION NUMBER: PCT/US95/07111

PRIOR FILING DATE: 1995-05-31

PRIOR APPLICATION NUMBER: US 08/250,856

PRIOR FILING DATE: 1994-05-31

NUMBER OF SEQ ID NOS: 109

SEQ ID NO 47

LENGTH: 20

OY 1 GTGCTCATTTGATGC 15

The strand cDNA was primed with a Not I - oligo(dA) primer
TGTACCAATCTGAAGTGGAAGCGCCGGTTTTTTT 3'.

double-stranded cDNA was size selected, ligated to Eco RI adapters (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of a modified pT73 vector (Pharmacia). Library constructed by Bento Soares and M. Fatima Bonaldo. "

ORIGIN

	Query Match	82.7%	Score 12.4	DB 1	Length 40
	Best Local Similarity	92.9%	Pred. No. 2e+04		
	Matches 13	Conservative	0	Mismatches 1	Indels 0
					Gaps 0
OY	2 TGTTCATTGATGC	15			
Ob	17 TCTTCATTGATGC	4			

RESULT 2

LOCUS	CL213712	41 bp	mRNA	linear	SS 30-JUN-2004
DEFINITION	A037F07 GTC Gene Trap Library GV03C04 Mus musculus CDNA clone				
ACCESSION	A037F07, mRNA sequence.				
VERSION	CL213712				
KEYWORDS	CL213712.2	GI:49489570			
SOURCE	GSS.				
ORGANISM	Mus musculus	(house mouse)			
	Mus musculus				

REFERENCE

AUTHORS	TITLE	JOURNAL
Innesen, J., Flores, T., Van Sloun, P., Fuchsdauer, B.M., Vauti, F., Arnold, H.H., Schultgen, P., Wurtz, W., Von Melchner, A and Rütz, P.	A large-scale, gene-driven mutagenesis approach for the functional analysis of the mouse genome	Proc.Natl. Acad. Sci. U.S.A. 100 (17), 9918-9922 (2003)

MEDLINE 22810117

PUBMED 12904583
COMMENT On Jun 30, 2004 this sequence version replaced gi:40730613

```
Email: info@genetrapp.de
prbetrageo gene trap. Sequence tag generated by 5'RACE. Additional
sequence information can be found at:
'http://genetrapp.gsf.de/project/web/new/database/result_clone.html?
clone_id=A037F07'. ES cell line harboring insertion mutation of
target gene is available at:
'http://genetrapp.gsf.de/project/web/new/order_clones/howtoorder.htm
l'. Inhouse Sequence Identifier: 09103
Class: Gene Trap.
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FEATURES

Source

```

/organism="Mus musculus"
/mol_type="mRNA"
/strain="129 Sv"
/db_xref="taxon:10090"
/clone="A037F07"
/sex="Male"
/cell_type="Embryonic stem cell"
/cell_line="ES cells 129S2 (formerly 129/SvPas)"
/clone_lib="GATC Gene Trap Library GV03C04"
/note="Vector: p1betaGeo"

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ORIGIN

	Query Match	Similarity	Score	DB	Length
Best Local	13	92.9%	92.9%	Pred. No. 2e+04	1
Matches	13	Conservative	0	Mismatches	1
				Indels	0
				Gaps	0
QY	2	TGCTCATTCGATGC	15		
db	21	TGCTCATTCGATGC	8		

RESULT 3

CW509288/C

LOCUS	CM509288	41 bp	mRNA	linear	GSS 06-OCT-2004
DEFINITION	BGA434 BayGenomics Gene Trap Library pGTL1xf Mus musculus cDNA, mRNA sequence.				
ACCESSION	CM509288				
VERSION	CM509288.1	GI:53838793			
KEYWORDS	GSS.				
SOURCE	Mus musculus (house mouse)				
ORGANISM	Mus musculus				
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognath; Muridae; Murinae; Mus. 1 (bases 1 to 41)				
AUTHORS	BayGenomics.				
TITLE	http://baygenomics.ucsf.edu/				
JOURNAL	Unpublished (2001)				
COMMENT	Contact: BayGenomics				

ORGANISM

REFERENCE	1 (bases 1 to 41)
AUTHORS	BayGenomics.
TITLE	http://baygenomics.ucsf.edu/
JOURNAL	Unpublished (2001)
COMMENT	Contact: BayGenomics

Sequence tag generated by 5' RAGE of total RNA from gene trap ES cell line. ES cell lines harboring insertion mutation of target gene are available upon request from BayGenomics. Annotation information available from http://baygenomics.ucsf.edu/cgi-bin/BaySearch.py?OPTION=EXACT&TYPE=CELL_LINE&KEY=BCGA434
Class: Gene Trap.

FEATURES

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source
1. .41
/organism="Mus musculus"
/mol_type="mRNA"
/strain="129 Ola"
/db_xref="taxon:10090"
/sex="Male"
/cell_type="Embryonic stem cell"
/clome_lib="BayGenomics Gene Trap Library pGTL1xf"
/clome_vector="pGTL1xf"

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ORIGIN

Query Match	82.7%	Score 12.4	DB 9	Length 41
Best Local Similarity	92.9%	Pred. No. 2e+4		
Matches 13	Conservative	0	Mismatches 1	Indels 0
Gaps				0
Qy	2	TGCTCATTCATGTC	15	
db	24	TGCTCATTCATGTC	11	

1

RESULT	4
CC894144/c	
LOCUS	CC894144
DEFINITION	CC894144 43 bp mRNA linear GSS 02-SEP-2003
RNAI38 BayGenomics Gene Trap Library pGT2LXf Mus musculus cdna,	
MRA sequence.	

ACCESSION

VERSION	CC894144.2	GI:34408991
KEYWORDS	GSS.	
SOURCE	Mus musculus	(house mouse)
ORGANISM	Mus musculus	

REFERENCE
AUTHORS
1 (bases 1 to 43)
BayGenomics.

TITLE

JOURNAL
Unpublished (2001)
COMMENT
On Sep 2, 2003 this sequence version replaced gi:33392557.

Email: info@baygenomics.ucsf.edu
Sequence tag generated by 5' RACE of total RNA from gene trap ES cell line. ES cell lines harboring insertion mutation of target gene are available upon request from Baygenomics. Annotation information available from http://baygenomics.ucsf.edu/cgi-bin/BaySearch.py?OPTION=EXACTYPE=CELL_LINETYPE=RRK138
Class: Gene Trap.

FEATURES

Location/Qualifiers

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SOURCE
1. 43
/organism="Mus musculus"
/mol_type="mRNA"
/strain="129 Ola"
/db_xref="taxon:10090"
/sex="Male"
/cell_type="Embryonic stem cell"
/clone_lib="BayGenomics Gene Trap Library pGT2Lxf"
/notes="Vector: pGT2Lxf"

ORIGIN
Query Match      82.7%  Score 12.4; DB 9; Length 43;
Best Local Similarity 92.9%  Pred. No. 2e+04;
Matches 13; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY
2 TGCTCCATTGATGC 15
|||||
26 TGCTCCATTGATGC 13

RESULT 5
CC200383      47 bp  mRNA  linear  GSS 09-MAY-2003
LOCUS
DEFINITION
RRE120 BayGenomics Gene Trap Library pGT2Lxf Mus musculus cDNA,
mRNA sequence.
ACCESSION
CC200383
VERSION
CC200383.1 GI:30480146
KEYWORDS
GSS.
SOURCE
Mus musculus (house mouse)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 47)
BayGenomics.
http://baygenomics.ucsf.edu/
Unpublished (2001)
Contact: BayGenomics
Bay Area Functional Genomics Consortium (BayGenomics)
Email: info@baygenomics.ucsf.edu
Sequence tag generated by 5' RACE of total RNA from gene trap ES
cell line. ES cell lines harboring insertion mutation of target
gene are available upon request from BayGenomics. Annotation
Information available from
http://baygenomics.ucsf.edu/cgi-bin/BaySearch.py?OPTION=EXACTTYPE=
CEL1.LINKEY=RRE120
Class: Gene Trap.
FEATURES
Location/Qualifiers
1..47
/organism="Mus musculus"
/mol_type="mRNA"
/strain="129 Ola"
/db_xref="taxon:10090"
/sex="Male"
/cell_type="Embryonic stem cell"
/clone_lib="BayGenomics Gene Trap Library pGT2Lxf"
/notes="Vector: pGT2Lxf"

ORIGIN
Query Match      82.7%  Score 12.4; DB 8; Length 47;
Best Local Similarity 92.9%  Pred. No. 2.1e+04;
Matches 13; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY
2 TGCTCCATTGATGC 15
|||||
30 TGCTCCATTGATGC 17

RESULT 6
CL659092      32 bp  DNA  linear  GSS 09-JUN-2004
LOCUS
DEFINITION
PRI0133a.G11 - PRI0133a.B21.1 (32) Mixed stage foemid library of P.
pacificus var. California Pristionchus pacificus genomic, genomic
survey sequence.

```

```

ACCESSION
CL659092
VERSION
CL659092.1 GI:50142802
KEYWORDS
GSS.
SOURCE
Pristionchus pacificus
ORGANISM
Pristionchus pacificus
Eukaryota; Metazoa; Nematoda; Chromadorea; Diplogasterida;
Neodiplogasteridae; Pristionchus.
1 (bases 1 to 32)
Strinvaasan,V., Otto,G.W., Kahlow,U., Geisler,R. and Sommer,R.J.
Appads: an Acedb database for the nematode satellite organism
Pristionchus pacificus
Nucleic Acids Res. 32 (1), D421-D422 (2004)
Contact: Sommer RJ
Evolutionary Biology
Max-Planck-Institute for Developmental Biology
Spemannstr. 37-39, Tuebingen D-72076, Germany
Tel: 00497071601371
Fax: 00497071601498
Email: ralf.sommer@tuebingen.mpg.de
This library was generated at Caltech, Pasadena, USA and end
sequenced at Vancouver, Canada.
Seq primer: T7
Class: fosmid ends.
FEATURES
Location/Qualifiers
1..32
/organism="Pristionchus pacificus"
/mol_type="genomic DNA"
/strain="California"
/db_xref="taxon:54126"
/clone_lib="Mixed stage foemid library of P. pacificus
var. California"
/notes="Vector: pBplfos-5 Fosmid vector"

ORIGIN
Query Match      78.7%  Score 11.8; DB 9; Length 32;
Best Local Similarity 86.7%  Pred. No. 4.4e+04;
Matches 13; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY
1 GTGCTCCATTGATGC 15
|||||
29 GTGCTCCATTGATGC 15

RESULT 7
AG190569      27 bp  DNA  linear  GSS 06-MAR-2004
LOCUS
DEFINITION
Pan troglodytes DNA, clone: RP43-066B21.TV, genomic survey
sequence.
ACCESSION
AG190569
VERSION
AG190569.1 GI:45222745
KEYWORDS
GSS.
SOURCE
Pan troglodytes (chimpanzee)
ORGANISM
Pan troglodytes
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Pan.
1
Park,H., Kim,Y., Kim,S., Han,Y., Woo,T., Park,K., Eun,C.J.,
Hoon,S.T., Chu,M., Kim,H., Joo,S., Kim,C., Song,W. and Yoo,H.
BAC end sequences of library RP-43
Unpublished
2 (bases 1 to 27)
Park,H., Kim,Y., Kim,S., Han,Y., Woo,T., Park,K., Eun,C.J.,
Hoon,S.T., Chu,M., Kim,H., Joo,S., Kim,C., Song,W. and Yoo,H.
Direct Submission
Submitted (07-JAN-2002) Hong-Seog Park, Korea Research Institute of
Bioscience and Biotechnology (KRIIB), Genome Research Center (GRC);
52, Oun-dong, Yuseong-gu, Daejeon 305-333, Korea
(E-mail:redetone@mail.kribb.re.kr, URL:http://pns.grc.kribb.re.kr/,
Tel:82-42-866-7181, Fax:82-42-860-4409)
Clones are derived from the chimpanzee BAC library RP-43 This BAC
end was generated during the R&D process and may have higher chance
of clone tracking errors.
PRIMERS
COMMENT

```

Sequencing: T7

LIBRARY : DBAc63.6
Vector : ECoRI
R.Site 1 : ECoRI
R.Site 2 : ECoRIFEATURES
source
Location/Qualifiers

1. .27
/organism="Pan troglodytes"
/mol_type="genomic DNA"
/db_xref="taxon:9598"
/clone="RP43-066B21.T7"
/sex="male"
/cell_type="lymphocytes"
/clone_lib="RP-43 Chimpanzee Male BAC library"

ORIGIN

Query Match 73.3%; Score 11; DB 9; Length 27;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GTGCTCCATTG 11
16 GTGCTCCATTG 26

RESULT 8 39 bp DNA linear GSS 13-DEC-2000
LOCUS A2579181
DEFINITION clone UUCGCM0365G10 F, genomic survey sequence.
ACCESSION A2579181
VERSION A2579181.1 GI:11693526
KEYWORDS GSS.

SOURCE
ORGANISM
Mus musculus (house mouse)

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 39)

REFERENCE
AUTHORS

Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamll,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T.,
Rilly,M., Rose,M., Rose,R., Stokes,R., Tinney,A., von
Niederhausern,A. and Wright,D.,Weiss,R.
Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts

JOURNAL
COMMENT

Unpublished (2000)
Contact: Robert B. Weiss
University of Utah Genome Center
Km. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0363 row: G column: 10
Seq primer: CGTGTAAACGACGCCACGT
Class: plasmid ends
High quality sequence stop: 39.
Location/Qualifiers

FEATURES
source

1. .39
/organism="Mus musculus"
/mol_type="genomic DNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUCGCM0365G10"
/sex="male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/clone_lib="Mouse 10kb plasmid UUCGCM library"
/note="Vector: pMD42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a

0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adapted DNA was purified and size-selected for a 9.5 to
10.5 Kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of pMD42 (g14732114|gb|AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptor complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
adapted vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

ORIGIN

Query Match 73.3%; Score 11; DB 8; Length 39;
Best Local Similarity 100.0%; Pred. No. 1.4e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 5 TCCATTGATGC 15
18 TCCATTGATGC 28

RESULT 9 46 bp mRNA linear GSS 21-SEP-2004
LOCUS CL982908
DEFINITION GC0134 TIGEM gene trap library Mus musculus cDNA clone 8644.24,
mRNA sequence.
ACCESSION CL982908
VERSION CL982908.1 GI:52420303
KEYWORDS GSS.

SOURCE
ORGANISM
Mus musculus (house mouse)

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 46)

REFERENCE
AUTHORS

Cobellis,G., Nicolaus,G., Marra,E., Barbarisi,M., Sardiello,M., Di
Giorgio,F.P., Iovino,N., Zollo,M., Ballabio,A. and Corsette,R.
Tagging genes with cassette-exchange sites
Unpublished (2004)
Contact: TIGEM
107

JOURNAL
COMMENT

Via P. Castellino, 111, 80131 NAPOLI, ITALY
Tel: +390815132205
Fax: +390815790919
Email: cobellis@tigem.it
Sequence tag generated by 5' RACE of total RNA from gene trap ES
cell line. ES cell lines harboring insertion mutation of target
gene are available upon request from TIGEM. Annotation information
available from TIGEM
Class: Gene Trap.

FEATURES
source

1. .46
/organism="Mus musculus"
/mol_type="mRNA"
/strain="129 Ola"
/db_xref="taxon:10090"
/clone="8644.24"
/sex="male"
/cell_type="Embryonic stem cell"
/cell_line="E14"
/clone_lib="TIGEM gene trap library"
/note="Vector: pFLIP1"

ORIGIN

Query Match 73.3%; Score 11; DB 9; Length 46;
Best Local Similarity 100.0%; Pred. No. 1.4e+05;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 CTCATTGATG 14

Db 26 CTCGATGATG 36

RESULT 10
LOCUS BX659267/c 49 bp DNA linear GSS 04-APR-2004

DEFINITION Arabidopsis thaliana T-DNA flanking sequence GK-644G02-022818,
genomic survey sequence.

ACCESSION BX659267

VERSION BX659267.1 GI:37615655

KEYWORDS GSS.

SOURCE Arabidopsis thaliana (chale crese)

ORGANISM

REFERENCE 1
AUTHORS Li,Y., Rosso,M.G., Strizhov,N., Viehoever,P. and Weishaar,B.
TITLE GABI-Kat Simplesearch: a flanking sequence tag (FST) database for the identification of T-DNA insertion mutants in Arabidopsis thaliana
JOURNAL Bioinformatics 19 (11), 1441-1442 (2003)
MEDLINE 22755829
PUBMED 12874060

REFERENCE 2
AUTHORS Rosso,M.G., Li,Y., Strizhov,N., Reis,B., Dekker,K. and Weishaar,B.
TITLE An Arabidopsis thaliana T-DNA mutagenized population (GABI-Kat) for flanking sequence tag-based reverse genetics
JOURNAL Plant Mol. Biol. 53 (1-2), 247-259 (2003)
MEDLINE 23117147
PUBMED 14756321

REFERENCE 3
AUTHORS Strizhov,N., Li,Y., Rosso,M.G., Viehoever,P., Dekker,K.A. and Weishaar,B.
TITLE High-throughput generation of sequence indexes from T-DNA mutagenized Arabidopsis thaliana lines
JOURNAL Biotechniques 35 (6), 1164-1168 (2003)
PUBMED 14682050

REFERENCE 4
AUTHORS Strizhov,N., Rosso,M.G., Li,Y. and Weishaar,B.
TITLE Direct Submission
Submitted (31-MAR-2004) Weishaar B., Max-Planck-Institut fuer Zuechungsforchung, Carl-von-Linne-Weg 10, Koeln, 50829, Germany
This sequence has been recovered from the left border of the T-DNA. It indicates an insertion close to or within gene Atlg72180.
Details on the protocols used for generation of the sequence are described in References 1-3. The sequences are generated at the MPI for Plant Breeding Research in the context of the GABI-Kat project.
GABI-Kat is part of the German Plant Genomics program designated 'GABI'. Information on line availability can be found at:
<http://www.mpiz-koeln.mpg.de/GABI-Kat/>.

FEATURES
source
1. 49
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="GK-644G02-022818"
/cclone_lib="Arabidopsis thaliana T-DNA insertion lines"
/ecotype="Col-0"
/note="PCR was performed on DNA from Arabidopsis thaliana plants (T1) which were transformed with the T-DNA from vector PAC161 (GenBank accession number: AF537514). The lines contain one or more T-DNA insertions. The DNA fragment(s) resulting from the PCR were directly sequenced to determine the genomic sequence flanking the insertion. T-DNA derived sequences were removed."

ORIGIN
Query Match 73.3%; Score 11; DB 9; Length 49;
Best Local Similarity 100.0%; Pred. No. 1.4e+05;

Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 4 CTCGATGATG 14
|||||
Db 44 CTCGATGATG 34

RESULT 11
LOCUS AZ769902 26 bp DNA linear GSS 16-FEB-2001

DEFINITION IM0571G06F Mouse 10kb plasmid UGCM library Mus musculus genomic clone UGCM0571G06 F, genomic survey sequence.

ACCESSION AZ769902

VERSION AZ769902.1 GI:12890529

KEYWORDS GSS.

SOURCE Mus musculus (house mouse)

ORGANISM

REFERENCE 1
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 26)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C., Islam,H., Longacre,S., Mahmood,M., Meenen,E., Pedersen,T., Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausen,A. and Wright,D.,Weiss,R.
Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
JOURNAL Unpublished (2000)
CONTACT: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: dunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0571 row: G column: 06
Seq primer: CGTGTAAACGACGCCAGT
Class: plasmid ends
High quality sequence stop: 26.

FEATURES
source
1. 26
/organism="Mus musculus"
/mol_type="genomic DNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/cclone="UGCM0571G06"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/cclone_lib="Mouse 10kb plasmid UGCM library"
/note="Vector: PWD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (<http://www.jax.org/resources/documents/dnares/>). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adapted DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pWD42 (g14732114|g147329072.1), a copy-number inducible derivative of plasmid RI. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adapted mouse DNA was annealed to adapted vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

ORIGIN
Query Match 72.0%; Score 10.8; DB 8; Length 26;
Best Local Similarity 85.7%; Pred. No. 1.7e+05;

Matches 12; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 GTGCTCATGTATG 14
 |||||
 12 GTGCTCCCTTGACG 25

RESULT 12
 AU256788/c 28 bp mRNA linear EST 25-APR-2002

LOCUS AU256788 3'-directed mouse cDNA library Mus musculus cDNA clone

DEFINITION BED0008978 3', mRNA sequence.

ACCESSION AU256788

VERSION AU256788

KEYWORDS EST.

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus (house mouse)

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

AUTHORS 1 (bases 1 to 28)

TITLE Kato, K. and Matoba, R.

JOURNAL Generation of expressed sequence tags from mouse brain

COMMENT Unpublished (2002)

Contact: Kikuya Kato
 Graduate School of Biological Sciences
 Nara Institute of Science and Technology
 8916-5 Takayama, Ikoma, Nara 630-0101, Japan
 Tel: 81-743-72-5581
 Fax: 81-743-72-5589
 Email: kkatob@aisc-nara.ac.jp,
 URL: http://love2.aisc-nara.ac.jp/BED/index.html.
 Location/Qualifiers

FEATURES
 source 1..28
 /organism="Mus musculus"
 /mol_type="mRNA"
 /db_xref="taxon:10090"
 /clone="BED0008978"
 /tissue_type="brain"
 /clone_lib="3'-directed mouse cDNA library"

ORIGIN

Query Match 72.0%; Score 10.8; DB 1; Length 28;
 Best Local Similarity 85.7%; Pred. No. 1.7e+05;
 Matches 12; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 2 TGCTCCATGTATG 15
 |||||
 19 TCCCTCATGTATG 6

RESULT 13
 AZ780454/c 28 bp DNA linear GSS 16-FEB-2001

LOCUS AZ780454 280017016R Mouse 10kb plasmid UGCG1M library Mus musculus genomic

DEFINITION clone UGCG2M0017016 R, genomic survey sequence.

ACCESSION AZ780454

VERSION AZ780454.1 GI:12912132

KEYWORDS GSS.

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus (house mouse)

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

AUTHORS 1 (bases 1 to 28)

TITLE Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C., Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly, M., Rose, W., Rose, R., Stokes, R., Tingey, A., von Niederhausen, A. and Wright, D. Weis, R.

JOURNAL Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts

COMMENT Unpublished (2000)
 Contact: Robert B. Weis
 University of Utah Genome Center

Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: dunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0017 row: 0 column: 16
 Seq primer: CACACAGAAACAGCTATGAC
 Class: plasmid ends
 High quality sequence stop: 28.

FEATURES
 source 1..28
 /organism="Mus musculus"
 /mol_type="genomic DNA"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UGCG2M0017016"
 /sex="Male"
 /lab_host="R. Coli strain XL10-Gold, T1-resistant, F-"
 /clone_lib="Mouse 10kb plasmid UGCG1M library"
 /note="Vector: pMD29v; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pMD42 (g14732114|gb|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

ORIGIN

Query Match 72.0%; Score 10.8; DB 8; Length 28;
 Best Local Similarity 85.7%; Pred. No. 1.7e+05;
 Matches 12; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 GTGCTCATGTATG 14
 |||||
 23 GTGCTCAGTTATG 10

RESULT 14
 TA59H06P/c 42 bp DNA linear GSS 13-DEC-2000

LOCUS TA59H06P T. brucei sheared genomic DNA clone 59h06, forward sequence,

DEFINITION genomic survey sequence.

ACCESSION AL455680

VERSION AL455680.1 GI:11857958

KEYWORDS GSS.

SOURCE Trypanosoma brucei

ORGANISM Trypanosoma brucei

REFERENCE Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae; Trypanosoma.

AUTHORS 1 (bases 1 to 42)

TITLE Hall, N., Bowman, S., Leonard, N.J., Doggett, J., Atkin, R., Chillingworth, C., Ormond, D., Harris, B., El-Sayed, N., Hou, L., Melville, S.B., Rajadream, M.A. and Barrell, B.G.

JOURNAL Direct Submission

COMMENT Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton, Cambridge CB10 1SA, E-mail: barrell@sanger.ac.uk and nh1@sanger.ac.uk
 Constructed at the Institute for Genomic Research (TIGR), Rockville, MD. Genomic DNA isolated from a cloned population of

Db 30 GTGCTCATGTAG 43
 Search completed: June 21, 2005, 06:51:11
 Job time : 3008 secs

Trypanosoma brucei (TREU927/4 GUTat 10.1) was mechanically sheared to give a tight size distribution (4 kb). The v + i method used for the library construction is described in detail in Smith, H. and Venter, J.C. (Making small insert libraries for whole genome shotgun sequencing projects. In Genome Sequencing: A Practical Approach, eds. M. Vaudin and B. Barrell, Oxford University Press, 1999).
 Email: neisayed@iclr.org
 Details of T. brucei sequencing at the Sanger Centre are available at http://www.sanger.ac.uk/Projects/T_brucei/.
 Location/Qualifiers

1. .42
 /organism="Trypanosoma brucei"
 /mol_type="genomic DNA"
 /strain="TREU927"
 /db_xref="taxon:5691"
 /clone="59h06"

ORIGIN

Query Match 72.0%; Score 10.8; DB 9; Length 42;
 Best Local Similarity 85.7%; Pred. No. 1.8e+05;
 Matches 12; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GTGCTCATGTAG 14
 |||||
 Db 22 GTGCCCCATTGAG 9

RESULT 15
 CV521218 43 bp mRNA linear EST 06-OCT-2004
 DEFINITION 0089P0056Z.x0_D02 Mimulus guttatus library 2 Mimulus guttatus cDNA
 LOCUS clone 0089P0056Z.x0_D02, mRNA sequence.
 CV521218
 ACCESSION CV521218
 VERSION CV521218.1 GI:53847750
 KEYWORDS EST.
 SOURCE Mimulus guttatus (spotted monkey flower)

ORGANISM Mimulus guttatus
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
 asterids; lamiales; Lamiales; incertae sedis; Mimuleae;
 Mimulus.

REFERENCE 1 (bases 1 to 43)
 AUTHORS Willis, J., Visions, T., Dietrich, F.S. and Allen, A.
 TITLE Mimulus guttatus cDNA sequence
 JOURNAL Unpublished (2004)
 COMMENT Contact: Willis J
 Department of Biology
 Duke University
 072-A Biological Sciences Science Drive, Durham, NC 27708, USA
 Tel: 919 660 7340
 Fax: 919 660 7293
 Email: jwillis@duke.edu
 Plate: 0089P0056 row: 02 column: D
 Seq primer: 77
 High quality sequence start: 80
 High quality sequence stop: 132.
 Location/Qualifiers

FEATURES
 source 1. .43
 /organism="Mimulus guttatus"
 /mol_type="mRNA"
 /db_xref="taxon:4155"
 /clone="0089P0056Z.x0_D02"
 /clone_1b="Mimulus guttatus library 2"
 /note="Vector: pGEM-T Easy; a Mimulus guttatus cDNA library"

ORIGIN

Query Match 72.0%; Score 10.8; DB 7; Length 43;
 Best Local Similarity 85.7%; Pred. No. 1.8e+05;
 Matches 12; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GTGCTCATGTAG 14

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: June 21, 2005, 01:08:35 ; Search time 3103 Seconds
(without alignments)
184.004 Million cell updates/sec

Title: US-10-075-994A-1
Perfect score: 15
Sequence: 1 gtgtccatgatgc 15

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database :

EST: *
1: gb_est1: *
2: gb_est2: *
3: gb_hlc: *
4: gb_est3: *
5: gb_est4: *
6: gb_est5: *
7: gb_est6: *
8: gb_ges1: *
9: gb_ges2: *

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	15	100.0	165	9	CG531606 OST114549
C 2	15	100.0	166	9	CG497924 OST18825
C 3	15	100.0	207	1	AA306005 EST177032
C 4	15	100.0	229	9	CG605299 OST81834
C 5	15	100.0	231	9	CG661765 OST43394
C 6	15	100.0	253	9	CG580338 OST19649
C 7	15	100.0	274	2	BF897689 QV1-MT022
C 8	15	100.0	276	9	CG664359 OST451337
C 9	15	100.0	282	9	CG546546 OST146106
C 10	15	100.0	293	2	BF468476 UI-M-CD0
C 11	15	100.0	294	1	AA332421 EST16381
C 12	15	100.0	300	9	CG577036 OST12398
C 13	15	100.0	300	9	CG591346 OST45811
C 14	15	100.0	308	9	CG513061 OST66238
C 15	15	100.0	309	7	CN295726 170005321
C 16	15	100.0	327	7	DA3526 DA3526 R1ce
C 17	15	100.0	328	5	BY329807 BY329807
C 18	15	100.0	335	5	BY324333 BY324333
C 19	15	100.0	340	5	BY068929 BY068929
C 20	15	100.0	346	5	BQ375054 WR4-TN011
C 21	15	100.0	349	5	BY063442 BY063442
C 22	15	100.0	351	9	CG633156 OST152509
C 23	15	100.0	353	6	CB779923 AMGNNUC:N
C 24	15	100.0	355	5	BY147345 BY147345

C 25	15	100.0	355	9	CG623442	CG623442 OST125479
C 26	15	100.0	358	5	BY014540	BY014540 BY014540
C 27	15	100.0	359	5	BY304066	BY304066 BY304066
C 28	15	100.0	362	5	BY036035	BY036035 BY036035
C 29	15	100.0	362	9	CG555213	CG555213 OST169619
C 30	15	100.0	366	5	BY063180	BY063180 BY063180
C 31	15	100.0	366	5	BY318790	BY318790 BY318790
C 32	15	100.0	367	5	BY014670	BY014670 BY014670
C 33	15	100.0	374	5	BY071821	BY071821 BY071821
C 34	15	100.0	376	5	BY280317	BY280317 BY280317
C 35	15	100.0	379	2	BE244097	BE244097 TCBAPIE06
C 36	15	100.0	379	5	BY047534	BY047534 BY047534
C 37	15	100.0	382	1	AU077313	AU077313 AU077313
C 38	15	100.0	384	5	BY087978	BY087978 BY087978
C 39	15	100.0	387	1	AA017736	AA017736 mh41b05.r
C 40	15	100.0	388	9	CG603416	CG603416 OST278044
C 41	15	100.0	391	5	BY061467	BY061467 BY061467
C 42	15	100.0	399	5	BY279803	BY279803 BY279803
C 43	15	100.0	400	5	BY288082	BY288082 BY288082
C 44	15	100.0	401	6	CB698634	CB698634 AMGNNUC:M
C 45	15	100.0	406	5	BY314742	BY314742 BY314742

ALIGNMENTS

RESULT 1
CG531606/c
LOCUS
DEFINITION
CG531606 165 bp mRNA linear GSS 01-OCT-2003
OST114549 Mus musculus 129Sv/Ev Mus musculus CDNA clone OST114549,
mRNA sequence.
ACCESSION
CG531606
VERSION
CG531606.1 GI:37318178
KEYWORDS
GSS.
SOURCE
Mus musculus (house mouse)
ORGANISM
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 165)
REFERENCE
Zambrowicz,B.P., Abuln,A., Ramirez-Solis,R., Richter,J.U.,
Piggott,J., Beltranderio,H., Buxton,E.C., Edwards,J., Finch,R.A.,
Fridde,C.J., Gupta,A., Hansen,G., Hu,Y., Huang,C.,
Key,B.W., Jr., Kipp,P., Kohlhauf,B., Ma,Z.-Q., Markesich,D.,
Payne,R., Potter,D.G., Qian,N., Shaw,J., Schrick,J., Shi,Z.-Z.,
Sparkes,M.J., Van Sligtenhorst,I., Vogel,P., Walke,W., Xu,N.,
Zhu,Q., Person,C. and Sands,A.T.
Wnt1 kinase deficiency lowers blood pressure in mice: a gene-trap
screen to identify potential targets for therapeutic intervention
Proc. Natl. Acad. Sci. U.S.A. 100 (24), 14109-14114 (2003)
Contact: Zambrowicz BP
OmniBank
Lexicon Genetics Incorporated
4000 Research Forest Drive, The Woodlands, TX 77381, USA
Email: materials@lexgen.com
Gene trap sequence tag generated by 3' RACE from mouse ES cells as
described in Zambrowicz et al (Nature, 1998 Apr 9;392(6676):608-11)
Class: Gene Trap.
FEATURES
source
1..165
/organism="Mus musculus"
/mol_type="mRNA"
/strain="129Sv/Ev"
/db_xref="taxon:10090"
/clone="OST114549"
/cell_type="embryonic stem cell"
/clone_lib="Mus musculus 129Sv/Ev"
ORIGIN
Query Match 100.0%; Score 15; DB 9; Length 165;
Best Local Similarity 100.0%; Pred. No. 7.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GTGTCCATTGATGC 15

Db 39 GTGCTCATGTATGC 25

|||||

RESULT 2
LOCUS CG497924/c
DEFINITION OST38825 Mus musculus 129Sv/Ev Mus musculus CDNA clone OST38825,
mRNA sequence.
ACCESSION CG497924
VERSION CG497924.1 GI:37267955
KEYWORDS GSS.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus

REFERENCE
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 166)
Zambrowicz,B.P., Abuin,A., Ramirez-Solis,R., Richter,L.J.,
Piggott,J., BeltrandeRio,H., Buxton,E.C., Edwards,J., Finch,R.A.,
Fridde,C.J., Gupta,A., Hansen,G., Hu,Y., Huang,W., Jaling,C.,
Key,B.W. Jr., Kipp,P., Kohlhauff,B., Ma,Z.-Q., Markesich,D.,
Payne,R., Potter,D.G., Qian,N., Shaw,J., Schrick,J., Shi,Z.-Z.,
Sparks,M.J., Van Sligtenhorst,I., Vogel,P., Walke,W., Xu,N.,
Zhu,Q., Person,C. and Sands,A.T.
Mki kinase deficiency lowers blood pressure in mice: a gene-trap
screen to identify potential targets for therapeutic intervention
Proc. Natl. Acad. Sci. U.S.A. 100 (24), 14109-14114 (2003)
Contact: Zambrowicz BP

TITLE
JOURNAL
COMMENT Omibank
Lexicon Genetics Incorporated
4000 Research Forest Drive, The Woodlands, TX 77381, USA
Email: materials@lexgen.com
Gene trap sequence tag generated by 3' RACE from mouse ES cells as
described in Zambrowicz et al (Nature. 1998 Apr 9;339(6676):608-11)
Class: Gene Trap.
Location/Qualifiers
1. 166
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/mol_type="mRNA"
/strain="129Sv/Ev"
/db_xref="taxon:10090"
/clone="OST38825"
/cell_type="embryonic stem cell"
/clone_1fb="Mus musculus 129Sv/Ev"

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Best Local Similarity 100.0%; Pred. No. 7.3e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GTGCTCATGTATGC 15
|||||

Db 35 GTGCTCATGTATGC 21
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RESULT 3
LOCUS AA306005
DEFINITION EST177032 Jurkat T-cells VI Homo sapiens CDNA 5' end similar to
Proto-oncogene rat mRNA sequence.
ACCESSION AA306005
VERSION AA306005.1 GI:1958375
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

REFERENCE
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
1 (bases 1 to 207)
Adams,M.D., Kerlavage,A.R., Fleischmann,R.D., Fuldner,R.A.,
Bull,C.J., Lee,N.H., Kirkness,E.F., Weinstock,K.G., Gocayne,J.D.,
White,O., Sutton,G., Blake,J.A., Brandon,R.C., Man-ai,C.,
Clayton,R.A., Cline,T.R., Cotton,M.D., Earle-Hughes,J., Fine,L.D.,
Fitzgerald,L.M., Fitzhugh,W.M., Fritchman,J.L., Geoghagen,N.S.,

Glodek,A., Gnehm,C.L., Hanna,M.C., Hedblom,E., Hinkle,P.S.Jr.,
Kelley,J.M., Kelley,J.C., Liu,L.-I., Marmaros,S.M., Merrick,J.M.,
Moreno-Palanges,R.F., McDonald,L.A., Nguyen,D.T., Pelligri,S.M.,
Phillips,C.A., Ryder,S.E., Scott,J.L., Saudet,D.M., Shirley,R.,
Small,K.V., Spriggs,T.A., Uteback,T.R., Weidman,J.F., Li,Y.,
Bednarek,D.P., Cao,L., Cepeda,M.A., Coleman,T.A., Collins,E.J.,
Dimke,D., Feng,D.-F., Ferrie,A., Fischer,C., Hastings,G.A.,
He,M.W., Hu,Y.S., Greene,J.M., Gruber,J., Hudson,P., Kim,A.K.,
Kozak,D.L., Kunsch,C., Hungjun,J., Li,H., Weisner,P.S., Olsen,H.,
Raymond,L., Wei,Y.F., Wing,J., Xu,C., Yu,G.L., Ruben,S.M.,
Dillon,P.J., Fannon,M.R., Rosen,C.A., Haseltine,W.A., Fields,C.,
Fraser,C.M. and Venter,J.C.
Initial assessment of human gene diversity and expression patterns
based upon 83 million nucleotides of CDNA sequence
Nature 377 (6547 Suppl), 3-174 (1995)
96026280

TITLE
JOURNAL
MEDLINE
PUBMED

COMMENT Other ESTs: TH0192945
Contact: Kerlavage, AR
Bioinformatics
The Institute for Genomic Research
9712 Medical Center Drive, Rockville, MD 20850 USA
Tel: 3018699056
Fax: 3018699423
Email: arkerlav@tigr.org
For clone availability, additional sequence and expression
information related to this EST, please check the TIGR Human Gene
Index (<http://www.tigr.org/cdb/hgi/hgi.html>)
Seq primer: M13 Reverse.
Location/Qualifiers
1. 207
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="FATCC (inhos):160136"
/db_xref="taxon:9606"
/cell_type="T-lymphocyte"
/clone_1fb="Jurkat T-cells VI"
/note="Vector: pBluescript SK-; Site_1: EcoRI; Site_2:
XhoI"

ORIGIN
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Best Local Similarity 100.0%; Pred. No. 7.6e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GTGCTCATGTATGC 15
|||||

Db 40 GTGCTCATGTATGC 26
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RESULT 4
LOCUS CG605299/c
DEFINITION OST281834 Mus musculus 129Sv/Ev Mus musculus CDNA clone OST281834,
mRNA sequence.
ACCESSION CG605299
VERSION CG605299.1 GI:37427977
KEYWORDS GSS.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus

REFERENCE
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 229)
Zambrowicz,B.P., Abuin,A., Ramirez-Solis,R., Richter,L.J.,
Piggott,J., BeltrandeRio,H., Buxton,E.C., Edwards,J., Finch,R.A.,
Fridde,C.J., Gupta,A., Hansen,G., Hu,Y., Huang,W., Jaling,C.,
Key,B.W. Jr., Kipp,P., Kohlhauff,B., Ma,Z.-Q., Markesich,D.,
Payne,R., Potter,D.G., Qian,N., Shaw,J., Schrick,J., Shi,Z.-Z.,
Sparks,M.J., Van Sligtenhorst,I., Vogel,P., Walke,W., Xu,N.,
Zhu,Q., Person,C. and Sands,A.T.
Mki kinase deficiency lowers blood pressure in mice: a gene-trap
screen to identify potential targets for therapeutic intervention
Proc. Natl. Acad. Sci. U.S.A. 100 (24), 14109-14114 (2003)

COMMENT Contact: Zambrowicz BP
OmiBank
Lexicon Genetics Incorporated
4000 Research Forest Drive, The Woodlands, TX 77381, USA
Email: material@lexgen.com
Gene trap sequence tag generated by 3' RACE from mouse ES cells as described in Zambrowicz et al (Nature. 1998 Apr 9;392(6676):608-11)
Class: Gene Trap.
Location/Qualifiers

FEATURES
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/mol_type="mRNA"
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/db_xref="taxon:10090"
/clone="OST281834"
/cell_type="embryonic stem cell"
/clone_id="Mus musculus 129Sv/Ev"

ORIGIN
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Best Local Similarity 100.0%; Pred. No. 7,7e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GTGCTCCATTGATGC 15
43 GTGCTCCATTGATGC 29

RESULT 5
CG661765/c 231 bp mRNA linear GSS 02-OCT-2003
LOCUS OST443394 Mus musculus 129Sv/Ev Mus musculus CDNA clone OST443394,
DEFINITION mRNA sequence.
ACCESSION CG661765
VERSION CG661765.1 GI:37485614
KEYWORDS GSS.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 231)
Zambrowicz, B.P., Abuin, A., Ramirez-Solis, R., Richer, L.J.,
Piggott, T., BeltrandelRio, H., Buxton, E.C., Edwards, J., Finch, R.A.,
Fridde, C.J., Gupta, A., Hansen, G., Hu, Y., Huang, W., Jaing, C.,
Key, B.W. Jr., Kipp, P., Kohlauf, B., Ma, Z.-Q., Markesich, D.,
Payne, R., Potter, D.G., Qian, N., Shaw, J., Schrick, J., Shi, Z.-Z.,
Spark, M.J., Van Sligtenhorst, I., Vogel, P., Walke, W., Xu, N.,
Zhu, Q., Person, C. and Sands, A.T.
Mki1 kinase deficiency lowers blood pressure in mice: a gene-trap
screen to identify potential targets for therapeutic intervention
Proc. Natl. Acad. Sci. U.S.A. 100 (24), 14109-14114 (2003)
Contact: Zambrowicz BP

JOURNAL OmiBank
COMMENT Lexicon Genetics Incorporated
4000 Research Forest Drive, The Woodlands, TX 77381, USA
Email: material@lexgen.com
Gene trap sequence tag generated by 3' RACE from mouse ES cells as described in Zambrowicz et al (Nature. 1998 Apr 9;392(6676):608-11)
Class: Gene Trap.
Location/Qualifiers

FEATURES
SOURCE 1..231
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/mol_type="mRNA"
/strain="129Sv/Ev"
/db_xref="taxon:10090"
/clone="OST443394"
/cell_type="embryonic stem cell"
/clone_id="Mus musculus 129Sv/Ev"

ORIGIN
Query Match 100.0%; Score 15; DB 9; Length 231;
Best Local Similarity 100.0%; Pred. No. 7,7e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GTGCTCCATTGATGC 15
30 GTGCTCCATTGATGC 16

RESULT 6
CG580338/c 253 bp mRNA linear GSS 02-OCT-2003
LOCUS OST219649 Mus musculus 129Sv/Ev Mus musculus CDNA clone OST219649,
DEFINITION mRNA sequence.
ACCESSION CG580338
VERSION CG580338.1 GI:37375289
KEYWORDS GSS.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 253)
Zambrowicz, B.P., Abuin, A., Ramirez-Solis, R., Richer, L.J.,
Piggott, T., BeltrandelRio, H., Buxton, E.C., Edwards, J., Finch, R.A.,
Fridde, C.J., Gupta, A., Hansen, G., Hu, Y., Huang, W., Jaing, C.,
Key, B.W. Jr., Kipp, P., Kohlauf, B., Ma, Z.-Q., Markesich, D.,
Payne, R., Potter, D.G., Qian, N., Shaw, J., Schrick, J., Shi, Z.-Z.,
Spark, M.J., Van Sligtenhorst, I., Vogel, P., Walke, W., Xu, N.,
Zhu, Q., Person, C. and Sands, A.T.
Mki1 kinase deficiency lowers blood pressure in mice: a gene-trap
screen to identify potential targets for therapeutic intervention
Proc. Natl. Acad. Sci. U.S.A. 100 (24), 14109-14114 (2003)
Contact: Zambrowicz BP

JOURNAL OmiBank
COMMENT Lexicon Genetics Incorporated
4000 Research Forest Drive, The Woodlands, TX 77381, USA
Email: material@lexgen.com
Gene trap sequence tag generated by 3' RACE from mouse ES cells as described in Zambrowicz et al (Nature. 1998 Apr 9;392(6676):608-11)
Class: Gene Trap.
Location/Qualifiers

FEATURES
SOURCE 1..253
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/clone="OST219649"
/cell_type="embryonic stem cell"
/clone_id="Mus musculus 129Sv/Ev"

ORIGIN
Query Match 100.0%; Score 15; DB 9; Length 253;
Best Local Similarity 100.0%; Pred. No. 7,9e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GTGCTCCATTGATGC 15
42 GTGCTCCATTGATGC 28

RESULT 7
BF897689 274 bp mRNA linear EST 18-JAN-2001
LOCUS QV1-MT0224-281100-512-b01 MT0224 Homo sapiens CDNA, mRNA sequence.
DEFINITION BF897689
ACCESSION BF897689
VERSION BF897689.1 GI:12289148
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE 1 (bases 1 to 274)
Dias Neto, E., Garcia Correa, R., Verjovski-Almeida, S., Briones, M.R.,
Nagai, M.A., da Silva, W. Jr., Zago, M.A., Bordin, S., Costa, F.P.,
Goldman, G.H., Carvalho, A.F., Matsumura, A., Bata, G.S., Simpson, D.H.,
Brunstein, A., deOliveira, P.S., Bucher, P., Jongeneel, C.V.,

O'Hare, M.J., Soares, F., Brentani, R.R., Reis, L.F., de Souza, S.J. and Simpson, A.J.
Shotgun sequencing of the human transcriptome with ORF expressed sequence tags
Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)

JOURNAL MEDLINE
20202663
10737800
Contact: Simpson A.J.G.
Laboratory of Cancer Genetics
Ludwig Institute for Cancer Research
Rua Prof. Antonio Frudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil
Tel: +55-11-2704922
Fax: +55-11-2707001
Email: asimpson@ludwig.org.br
This sequence was derived from the FAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL
(http://www.ludwig.org.br/scripts/gethtml2.pl?l=QV1&t2=QV1-MT0224-281100-512-D01&t3=2000-11-28&t4=1)
Seq primer: puc 18 forward
High quality sequence start: 27
High quality sequence stop: 274.
Location/Qualifiers

FEATURES
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/mol_type="mRNA"
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/clone_lib="MT0224"
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ORIGIN
Query Match 100.0%; Score 15; DB 2; Length 274;
Best Local Similarity 100.0%; Pred. No. 8e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
1 GTGCTCCATTGATGC 15
170 GTGCTCCATTGATGC 184

Db

RESULT 8
CG664359/c
LOCUS
DEFINITION
CG664359
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
COMMENT

CG664359 276 bp mRNA linear GSS 02-OCT-2003
OST451337 Mus musculus 129Sv/Ev Mus musculus cDNA clone OST451337,
mRNA sequence.
CG664359
CG664359.1 GI:37488208
GSS.
Mus musculus (house mouse)
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 276)
Zambrowicz, B.P., Abuin, A., Ramirez-Solis, R., Richter, L.J.,
Piggott, J., Beltranderio, H., Buxton, E.C., Edwards, J., Finch, R.A.,
Fridde, C.J., Gupta, A., Hansen, G., Hu, Y., Huang, W., Jaling, C.,
Key, B.W. Jr., Kipp, P., Kohlhauff, B., Ma, Z.-Q., Markesich, D.,
Payne, R., Potter, D.G., Qian, N., Shaw, J., Schrick, J., Shi, Z.-Z.,
Sparks, M.J., Van Sligtenhorst, I., Vogel, P., Walke, W., Xu, N.,
Zhu, Q., Person, C. and Sands, A.T.
Wnk1 kinase deficiency lowers blood pressure in mice: a gene-trap
screen to identify potential targets for therapeutic intervention
Proc. Natl. Acad. Sci. U.S.A. 100 (24), 14109-14114 (2003)
Contact: Zambrowicz BP
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Lexicon Genetics Incorporated

4000 Research Forest Drive, The Woodlands, TX 77381, USA
Email: materials@lexgen.com
Gene trap sequence tag generated by 3' RACE from mouse ES cells as
described in Zambrowicz et al (Nature. 1998 Apr 9;392(6676):608-11)
Class: Gene trap.
Location/Qualifiers

FEATURES
Source
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/cell_type="embryonic stem cell"
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ORIGIN
Query Match 100.0%; Score 15; DB 9; Length 282;
Best Local Similarity 100.0%; Pred. No. 8e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
1 GTGCTCCATTGATGC 15
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Db

RESULT 9
CG546546/c
LOCUS
DEFINITION
CG546546
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
COMMENT

CG546546 282 bp mRNA linear GSS 01-OCT-2003
OST16106 Mus musculus 129Sv/Ev Mus musculus cDNA clone OST16106,
mRNA sequence.
CG546546
CG546546.1 GI:37333133
GSS.
Mus musculus (house mouse)
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 282)
Zambrowicz, B.P., Abuin, A., Ramirez-Solis, R., Richter, L.J.,
Piggott, J., Beltranderio, H., Buxton, E.C., Edwards, J., Finch, R.A.,
Fridde, C.J., Gupta, A., Hansen, G., Hu, Y., Huang, W., Jaling, C.,
Key, B.W. Jr., Kipp, P., Kohlhauff, B., Ma, Z.-Q., Markesich, D.,
Payne, R., Potter, D.G., Qian, N., Shaw, J., Schrick, J., Shi, Z.-Z.,
Sparks, M.J., Van Sligtenhorst, I., Vogel, P., Walke, W., Xu, N.,
Zhu, Q., Person, C. and Sands, A.T.
Wnk1 kinase deficiency lowers blood pressure in mice: a gene-trap
screen to identify potential targets for therapeutic intervention
Proc. Natl. Acad. Sci. U.S.A. 100 (24), 14109-14114 (2003)
Contact: Zambrowicz BP
Omnibank
Lexicon Genetics Incorporated
4000 Research Forest Drive, The Woodlands, TX 77381, USA
Email: materials@lexgen.com
Gene trap sequence tag generated by 3' RACE from mouse ES cells as
described in Zambrowicz et al (Nature. 1998 Apr 9;392(6676):608-11)
Class: Gene trap.
Location/Qualifiers

FEATURES
Source
1. 282
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ORIGIN
Query Match 100.0%; Score 15; DB 9; Length 282;
Best Local Similarity 100.0%; Pred. No. 8e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
1 GTGCTCCATTGATGC 15
|||||

	EST.	Homo sapiens (human)		
SOURCE		Homo sapiens		
ORGANISM		Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo. 1 (bases 1 to 294) Adams,M.D., Kerlavage,A.R., Fleischmann,R.D., Fuldner,R.A., Bult,C.J., Lee,N.H., Kirkness,E.F., Weinstock,K.G., Gocayne,J.D., White,O., Sutton,G., Blake,J.A., Brandon,R.C., Man-ai,C., Clayton,R.A., Cline,T.R., Cotton,M.D., Earle-Hughes,J., Fine,L.D., Fitzgerald,L.M., Flitghun,W.M., Frithman,J.L., Georgagen,N.S., Gloder,A., Gnesh,C.L., Hanna,M.C., Hedblom,E., Hinkle,P.S Jr., Kellely,J.M., Kellely,J.C., Liu,L.-I., Marmaros,S.M., Merrick,J.M., Moreno-Palanguas,R.F., McDonald,L.A., Nguyen,D.T., Pelligrino,S.M., Phillips,C.A., Ryder,S.E., Scott,J.U., Sauder,D.W., Shirley,R., Snaili,K.V., Spriggs,T.A., Utepbach,T.R., Weidman,J.P., Li,Y., Bednariak,D.P., Cao,L., Cepeda,M.A., Coleman,T.A., Collins,L.Y., Dinke,D., Peng,D.-F., Ferrie,A.C., Fischer,C., Hastings,G.A., He,M.W., Hu,J.S., Greene,J.M., Gruber,J., Hudson,P., Kim,A.K., Kozak,D.L., Kunisch,C., Hungjun,U., Li,H., Meltsner,P.S., Olsen,H., Raymond,U., Wei,Y.F., Wing,J., Xu,C., Yu,G.L., Ruben,S.M., Dillon,P.J., Fannon,M.R., Rosen,C.A., Haseltine,W.A., Fields,C., Fraser,C.M. and Venter,J.C. Initial assessment of human gene diversity and expression patterns based upon 83 million nucleotides of cDNA sequence Nature 377 (6547 Suppl), 3-174 (1995) 96026280		
JOURNAL MEDLINE PUBMED COMMENT		Other ESTs: THC192945 Contact: Kerlavage, AR Bioinformatics The Institute for Genomic Research 9712 Medical Center Drive, Rockville, MD 20850 USA Tel.: 3018699056 Fax: 3018699423 Email: arkerlav@tigr.org For clone availability, additional sequence and expression information related to this EST, please check the TIGR Human Gene Index (http://www.tigr.org/tgr/hgi/hgi.html) Seq primer: M13 Reverse.		
FEATURES		Location/Qualifiers 1..294 /organism="Homo sapiens" /mol_type="mRNA" /db_xref="ATCC (inhost):134082" /db_xref="taxon:9606" /dev_stage="embryo, 8 wks" /clone_id="Embryo, 8 week I" /note="Organ: Embryo, 8 weeks; Vector: pBluescript SK-; Site_1: EcoRI; Site_2: XhoI"		
ORIGIN				
Query Match	100.0%; Score 15;	DB 1:	Length 294;	
Best Local Similarity	100.0%; Prid. No. 8.1e+02;			
Matches	15; Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;
Oy	1 GTGCTCATGTATGC	15		
Db	40 GTGCTCAATTATTC	26		
RESULT 12 CG577036/c LOCUS DEFINITION ACCESSION VERSION KEYWORDS SOURCE ORGANISM				
CG577036 OS2121398 Mus musculus 12SV/Ev Mus musculus CDNA clone OS2121398, mRNA Sequence. CG577036 CG577036.1 GI:37367640 GSS.				
Mus musculus (house mouse) Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.				

```

REFERENCE
AUTHORS      1 (bases 1 to 300)
              Zambrowicz,B.P., Abulin,A., Ramirez-Solis,R., Richter,L.J.,
              Piggett,J., BeltranderRio,H., Buxton,E.C., Edwards,J., Finch,R.A.,
              Fiddle,C.J., Gupta,A., Hansen,G., Hu,Y., Huang,W., Jaing,C.,
              Key,B.W., Jr., Klipp,P., Kohlhauff,B., Ma,Z.-Q., Markesich,D.,
              Payne,R., Potter,D.G., Qian,N., Shaw,J., Schrick,J., Shi,Z.-Z.,
              Sparks,M.J., Van Sligtenhorst,I., Vogel,P., Walke,W., Xu,N.,
              Zhu,Q., Person,C. and Sands,A.T.
TITLE        Wnt1 kinase deficiency lowers blood pressure in mice: a gene-trap
JOURNAL      screen to identify potential targets for therapeutic intervention
COMMENT      Proc. Natl. Acad. Sci. U.S.A. 100 (24), 14109-14114 (2003)
              Contact: Zambrowicz BP
              OmniBank
              Lexicon Genetics Incorporated
              4000 Research Forest Drive, The Woodlands, TX 77381, USA
              Email: materials@lexgen.com
              Gene trap sequence tag generated by 3' RACE from mouse ES cells as
              described in Zambrowicz et al (Nature. 1998 Apr 9;392(6676):608-11)
              Class: Gene Trap.

FEATURES
source       Location/Qualifiers
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              /cell_type="embryonic stem cell"
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ORIGIN
Query Match      100.0%; Score 15; DB 9; Length 300;
Best Local Similarity 100.0%; Pred. No. 8.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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DEFINITION OST245811 Mus musculus 129Sv/Ev Mus musculus cDNA clone OST245811,
            mRNA sequence.
ACCESSION  CG591346
VERSION    CG591346.1 GI:37399162
KEYWORDS   GSS.
SOURCE     Mus musculus (house mouse)
ORGANISM   Mus musculus
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
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REFERENCE
AUTHORS      1 (bases 1 to 300)
              Zambrowicz,B.P., Abulin,A., Ramirez-Solis,R., Richter,L.J.,
              Piggett,J., BeltranderRio,H., Buxton,E.C., Edwards,J., Finch,R.A.,
              Fiddle,C.J., Gupta,A., Hansen,G., Hu,Y., Huang,W., Jaing,C.,
              Key,B.W., Jr., Klipp,P., Kohlhauff,B., Ma,Z.-Q., Markesich,D.,
              Payne,R., Potter,D.G., Qian,N., Shaw,J., Schrick,J., Shi,Z.-Z.,
              Sparks,M.J., Van Sligtenhorst,I., Vogel,P., Walke,W., Xu,N.,
              Zhu,Q., Person,C. and Sands,A.T.
TITLE        Wnt1 kinase deficiency lowers blood pressure in mice: a gene-trap
JOURNAL      screen to identify potential targets for therapeutic intervention
COMMENT      Proc. Natl. Acad. Sci. U.S.A. 100 (24), 14109-14114 (2003)
              Contact: Zambrowicz BP
              OmniBank
              Lexicon Genetics Incorporated
              4000 Research Forest Drive, The Woodlands, TX 77381, USA
              Email: materials@lexgen.com
              Gene trap sequence tag generated by 3' RACE from mouse ES cells as
              described in Zambrowicz et al (Nature. 1998 Apr 9;392(6676):608-11)
              Class: Gene Trap.

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RESULT 15
LOCUS      CN295726      309 bp      mRNA      linear      EST 16-MAY-2004
DEFINITION 17000532192143 GRN_ES Homo sapiens cDNA 5', mRNA sequence.
ACCESSION  CN295726

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REFERENCE
AUTHORS      1 (bases 1 to 308)
              Zambrowicz,B.P., Abulin,A., Ramirez-Solis,R., Richter,L.J.,
              Piggett,J., BeltranderRio,H., Buxton,E.C., Edwards,J., Finch,R.A.,
              Fiddle,C.J., Gupta,A., Hansen,G., Hu,Y., Huang,W., Jaing,C.,
              Key,B.W., Jr., Klipp,P., Kohlhauff,B., Ma,Z.-Q., Markesich,D.,
              Payne,R., Potter,D.G., Qian,N., Shaw,J., Schrick,J., Shi,Z.-Z.,
              Sparks,M.J., Van Sligtenhorst,I., Vogel,P., Walke,W., Xu,N.,
              Zhu,Q., Person,C. and Sands,A.T.
TITLE        Wnt1 kinase deficiency lowers blood pressure in mice: a gene-trap
JOURNAL      screen to identify potential targets for therapeutic intervention
COMMENT      Proc. Natl. Acad. Sci. U.S.A. 100 (24), 14109-14114 (2003)
              Contact: Zambrowicz BP
              OmniBank
              Lexicon Genetics Incorporated
              4000 Research Forest Drive, The Woodlands, TX 77381, USA
              Email: materials@lexgen.com
              Gene trap sequence tag generated by 3' RACE from mouse ES cells as
              described in Zambrowicz et al (Nature. 1998 Apr 9;392(6676):608-11)
              Class: Gene Trap.

FEATURES
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              /clone_1ib="Mus musculus 129Sv/Ev"

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Best Local Similarity 100.0%; Pred. No. 8.2e+02;
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QY              1 GTGCTCATTGATGC 15
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RESULT 15
LOCUS      CN295726      309 bp      mRNA      linear      EST 16-MAY-2004
DEFINITION 17000532192143 GRN_ES Homo sapiens cDNA 5', mRNA sequence.
ACCESSION  CN295726

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VERSION CN295726.1 GI:47312140
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
AUTHORS 1 (bases 1 to 309)
Brandenberger,R., Wei,H., Zhang,S., Lei,S., Murage,J., Fisk,G.J.,
Li,Y., Xu,C., Fang,R., Guegler,K., Rao,M.S., Mandalam,R.,
Lebkowski,J and Stanton,L.W.
TITLE Transcriptional characterization elucidates signaling networks that
control human ES cell growth and differentiation
JOURNAL Nat. Biotechnol. 22 (6), 707-716 (2004)
COMMENT Contact: Brandenberger R
Regenerative Medicine
Geron Corporation
230 Constitution Drive, Menlo Park, CA 94025, USA
Tel: 650 473 8658
Fax: 650 473 7760
Email: rbrandenberger@geron.com
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H9"
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Search completed: June 21, 2005, 03:49:24
Job time : 3110 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: June 20, 2005, 20:52:15 ; Search time 1678 Seconds
(without alignments)
433.152 Million cell updates/sec

Title: US-10-075-994A-1
Sequence: 1 gtgctccattgatgc 15

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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2: gb_hlg:*
3: gb_in:*
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14: gb_vi:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	length	DB ID	Description
1	15	100.0	15	6	AR110775 Sequence
2	15	100.0	15	6	AR110777 Sequence
3	15	100.0	15	6	AR167449 Sequence
4	15	100.0	15	6	CQ789712 Sequence
5	15	100.0	15	6	AR310685 Sequence
6	15	100.0	15	6	AR310687 Sequence
7	15	100.0	15	6	AX797662 Sequence
8	15	100.0	15	6	AX957646 Sequence
9	15	100.0	15	6	AX957740 Sequence
10	15	100.0	15	6	AX958145 Sequence
11	15	100.0	15	6	BD106498 Sequence
12	15	100.0	20	6	AR073978 Sequence
13	15	100.0	20	6	AR216002 Sequence
14	15	100.0	25	6	AR110776 Sequence
15	15	100.0	25	6	AR310686 Sequence
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17	15	100.0	1496	8	AK064133 Oryza sat
18	15	100.0	2524	10	RA7RAFA M55427 Rat c-raf p
19	15	100.0	2811	10	AB057663 Mus muscu

C 20	15	100.0	2935	10	BC015273	BC015273 Mus muscu
C 21	15	100.0	2970	10	BC062071	BC062071 Rattus no
C 22	15	100.0	2975	6	AR473559	AR473559 Sequence
C 23	15	100.0	2977	6	AR073995	AR073995 Sequence
C 24	15	100.0	2977	6	AR110473	AR110473 Sequence
C 25	15	100.0	2977	6	BD237320	BD237320 Modulatio
C 26	15	100.0	2977	6	134402	134402 Sequence 17
C 27	15	100.0	2977	6	196180	196180 Sequence 17
C 28	15	100.0	2977	6	AR215978	AR215978 Sequence
C 29	15	100.0	2977	6	AR337749	AR337749 Sequence
C 30	15	100.0	2977	6	AR360020	AR360020 Sequence
C 31	15	100.0	2977	6	AX022819	AX022819 Sequence
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C 35	15	100.0	2977	6	AX696362	AX696362 Sequence
C 36	15	100.0	2977	6	AX777762	AX777762 Sequence
C 37	15	100.0	2977	6	BD091455	BD091455 Transgeni
C 38	15	100.0	2977	9	HSRAFR	X03484 Human mRNA
C 39	15	100.0	2981	6	CO723686	CO723686 Sequence
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C 42	15	100.0	71328	2	AC025497	AC025497 Homo sapi
C 43	15	100.0	82374	9	AY271661	AY271661 Homo sapi
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ALIGNMENTS

RESULT 1
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DEFINITION AR110775
ACCESSION AR110775
VERSION AR110775.1 GI:12827623
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 15)
AUTHORS Kasid,U., Gokhale,P., Dritschilo,A. and Rahman,A.
TITLES Liposomes containing oligonucleotides
JOURNAL Patent: US 6126965-A 1 03-OCT-2000;
FEATURES
LOCATION/Qualifiers
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source
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ORIGIN

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Best Local Similarity 100.0%; Pred. No. 1.2e+03;
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C 24 15 100.0 2977 6 AR110473
C 25 15 100.0 2977 6 BD237320
C 26 15 100.0 2977 6 134402
C 27 15 100.0 2977 6 196180
C 28 15 100.0 2977 6 AR215978
C 29 15 100.0 2977 6 AR337749
C 30 15 100.0 2977 6 AR360020
C 31 15 100.0 2977 6 AX022819
C 32 15 100.0 2977 6 AX030539
C 33 15 100.0 2977 6 AX337827
C 34 15 100.0 2977 6 AX622838
C 35 15 100.0 2977 6 AX696362
C 36 15 100.0 2977 6 AX777762
C 37 15 100.0 2977 6 BD091455
C 38 15 100.0 2977 9 HSRAFR
C 39 15 100.0 2981 6 CO723686
C 40 15 100.0 3216 9 BC018119
C 41 15 100.0 66275 2 AC100586
C 42 15 100.0 71328 2 AC025497
C 43 15 100.0 82374 9 AY271661
C 44 15 100.0 93614 9 AL139294
C 45 15 100.0 112234 8 AP006147
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LOCUS AR110777/c 15 bp DNA linear PAT 14-FEB-2001
DEFINITION Sequence 3 from patent US 6126965.
ACCESSION AR110777
VERSION AR110777.1 GI:12827625
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 15)
AUTHORS Kasid,U., Gokhale,P., Dritschilo,A. and Rahman,A.
TITLES Liposomes containing oligonucleotides

JOURNAL Patent: US 6126965-A 3 03-OCT-2000;
FEATURES Location/Qualifiers
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Db 15 GTGCTCCATTGATGC 1

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LOCUS Sequence 15 from patent US 6287591.
DEFINITION AR167449
ACCESSION AR167449
VERSION AR167449.1 GI:17903229
KEYWORDS
SOURCE
ORGANISM
REFERENCE 1 (bases 1 to 15)
AUTHORS Sample,S.C., Kilmuk,S.K., Harasym,T., Hope,M.J., Ansell,S.M.,
Cullis,P., Scherrer,P. and Debeyer,D.
TITLE Charged therapeutic agents encapsulated in lipid particles
JOURNAL Patent: US 6287591-A 15 11-SEP-2001;
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LOCUS Sequence 1 from Patent WO2004017944.
DEFINITION CQ789712
ACCESSION CQ789712
VERSION CQ789712.1 GI:45823264
KEYWORDS
SOURCE
ORGANISM
REFERENCE 1 (bases 1 to 15)
AUTHORS Zhang,J.A. and Ahmad,I.
TITLE Liposomal gentamicin compositions for better drug delivery
JOURNAL Patent: WO.2004017944-A 1 04-MAR-2004;
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DEFINITION AR310685
ACCESSION AR310685
VERSION AR310685.1 GI:31703829
KEYWORDS
SOURCE
ORGANISM
REFERENCE 1 (bases 1 to 15)
AUTHORS Kasid,U., Gokhale,P., Zhang,C., Dritschilo,A. and Rahman,A.
TITLE Cationic liposomal delivery system and therapeutic use thereof
JOURNAL Patent: US 6559129-A 1 06-MAY-2003;
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Db 1 GTGCTCCATTGATGC 15

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DEFINITION AR310687
ACCESSION AR310687
VERSION AR310687.1 GI:31703831
KEYWORDS
SOURCE
ORGANISM
REFERENCE 1 (bases 1 to 15)
AUTHORS Kasid,U., Gokhale,P., Zhang,C., Dritschilo,A. and Rahman,A.
TITLE Cationic liposomal delivery system and therapeutic use thereof
JOURNAL Patent: US 6559129-A 3 06-MAY-2003;
FEATURES Location/Qualifiers
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LOCUS Sequence 25 from Patent WO03039595.
DEFINITION AX797662
ACCESSION AX797662
VERSION AX797662.1 GI:37518090
KEYWORDS
SOURCE
ORGANISM Homo sapiens (human)

REFERENCE
AUTHORS
TITLE
JOURNAL
FEATURES
SOURCE

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
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Semple,S., Klimuk,S. and Yuan,Z.N.
Mucosal adjuvants comprising an oligonucleotide and a cationic
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Patent: WO 03039595-A 25 15-MAY-2003;
Inex Pharmaceuticals Corp. (CA)
Location/Qualifiers
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Db 1 GTGCTCCATTGATGC 15

RESULT 8
AX957646 15 bp DNA linear PAT 08-JAN-2004

LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

AX957646 Sequence 25 from Patent WO03094963.
AX957646
AX957646.1 GI:40785518
Homo sapiens (human)
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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
1
Tan,Y.K., Semple,S., Klimuk,S. and Chikh,G.
Methylated immunostimulatory oligonucleotides and methods of using
the same
Patent: WO 03094963-A 25 20-NOV-2003;
Inex Pharmaceuticals Corporation (CA)
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FEATURES
SOURCE

JOURNAL
Inex Pharmaceuticals Corporation (CA)

ORIGIN

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LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

AX957740 Sequence 25 from Patent WO03094828.
AX957740
AX957740.1 GI:40785558
Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
1
Tan,Y.K., Semple,S., Klimuk,S. and Chikh,G.
Cancer vaccines and methods of using the same
Patent: WO 03094828-A 25 20-NOV-2003;
Inex Pharmaceuticals Corp. (CA)

REFERENCE
AUTHORS
TITLE
JOURNAL

FEATURES
SOURCE

Location/Qualifiers
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OY 1 GTGCTCCATTGATGC 15
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Db 1 GTGCTCCATTGATGC 15

RESULT 10
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LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

AX958145 Sequence 25 from Patent WO03094829.
AX958145
AX958145.1 GI:40785809
Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
1
Semple,S., Chikh,G., Hope,M.J. and Tam,Y.K.
Pathogen vaccines and methods for using the same
Patent: WO 03094829-A 25 20-NOV-2003;
Inex Pharmaceuticals Corp. (CA)
Location/Qualifiers
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/organism="Homo sapiens"
/mol_type="unassigned DNA"
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FEATURES
SOURCE

JOURNAL
Inex Pharmaceuticals Corp. (CA)

ORIGIN

Query Match 100.0%; Score 15; DB 6; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.2e+03;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 1 GTGCTCCATTGATGC 15

RESULT 11
BD106498 15 bp DNA linear PAT 18-SEP-2002

LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

BD106498 High efficiency encapsulation of charged therapeutic agents in
lipid vesicles.
BD106498
BD106498.1 GI:23201316
JP 2002501511-A/15.
Chlamydia sp.
Chlamydia sp.
Bacteria; Chlamydiae; Chlamydiales; Chlamydiaceae; Chlamydia.
1 (bases 1 to 15)
Semple,S.C., Klimuk,S.K., Harasym,T., Hope,M.J., Ansel,S.M.,
Cullis,P., Scherret,P. and Debever,D.S.
High efficiency encapsulation of charged therapeutic agents in
lipid vesicles
Patent: JP 2002501511-A 15 15-JAN-2002;
INEX PHARMACEUTICALS CORP
PN JP 2002501511-A/15

JOURNAL
INEX PHARMACEUTICALS CORP

COMMENT

PD 15-JAN-2002
PF 14-MAY-1998 UP 1998548646
PI SEAN C SEMPLE,SANDRA K KLIMUK,TROY HARASYM,MICHAEL J HOPE, PI
STEVEN M ANSELL,
PI PETER CULLIS,PETER SCHERRER,DAN SUITE DBEYER PC AGIK9/00
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CC Topology: Linear; Location/Qualifiers.
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Db
 1 GTGCTCCATTGATGC 15
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LOCUS
DEFINITION Sequence 47 from patent US 5952229.
ACCESSION AR073978
VERSION AR073978.1 GI:10000738
KEYWORDS
SOURCE
ORGANISM Unknown.
REFERENCE
 AUTHORS Monia,B.P. and Boggs,R.T.
 TITLE Antisense oligonucleotide modulation of raf gene expression
 JOURNAL Patent: US 5952229-A 47 14-SEP-1999;
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ORIGIN
 Query Match
 100.0%; Score 15; DB 6; Length 20;
 Best Local Similarity 100.0%; Pred. No. 1.1e+03;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY
 1 GTGCTCCATTGATGC 15
 5 GTGCTCCATTGATGC 19
Db
 5 GTGCTCCATTGATGC 19
RESULT 13
AR216002
LOCUS
DEFINITION Sequence 49 from patent US 6410518.
ACCESSION AR216002
VERSION AR216002.1 GI:23314290
KEYWORDS
SOURCE
ORGANISM Unknown.
REFERENCE
 AUTHORS Monia,B.P.
 TITLE Antisense oligonucleotide inhibition of raf gene expression
 JOURNAL Patent: US 6410518-A 49 25-JUN-2002;
 FEATURES Location/Qualifiers
 1..20
 /organism="unknown"
 /mol_type="genomic DNA"
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 100.0%; Score 15; DB 6; Length 20;
 Best Local Similarity 100.0%; Pred. No. 1.1e+03;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY
 1 GTGCTCCATTGATGC 15
 |||||

Db
 5 GTGCTCCATTGATGC 19
RESULT 14
AR110776
LOCUS
DEFINITION Sequence 2 from patent US 6126965.
ACCESSION AR110776
VERSION AR110776.1 GI:12827624
KEYWORDS
SOURCE
ORGANISM Unknown.
REFERENCE
 AUTHORS Kasid,U., Gokhale,P., Dritschilo,A. and Rahman,A.
 TITLE Liposomes containing oligonucleotides
 JOURNAL Patent: US 6126965-A 2 03-OCT-2000;
 FEATURES Location/Qualifiers
 1..25
 /organism="unknown"
 /mol_type="unassigned DNA"
ORIGIN
 Query Match
 100.0%; Score 15; DB 6; Length 25;
 Best Local Similarity 100.0%; Pred. No. 1.1e+03;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY
 1 GTGCTCCATTGATGC 15
 8 GTGCTCCATTGATGC 22
Db
 8 GTGCTCCATTGATGC 22
RESULT 15
AR310686
LOCUS
DEFINITION Sequence 2 from patent US 6559129.
ACCESSION AR310686
VERSION AR310686.1 GI:31703830
KEYWORDS
SOURCE
ORGANISM Unknown.
REFERENCE
 AUTHORS Kasid,U., Gokhale,P., Zhang,C., Dritschilo,A. and Rahman,A.
 TITLE Cationic liposomal delivery system and therapeutic use thereof
 JOURNAL Patent: US 6559129-A 2 06-MAY-2003;
 FEATURES Location/Qualifiers
 1..25
 /organism="unknown"
 /mol_type="genomic DNA"
ORIGIN
 Query Match
 100.0%; Score 15; DB 6; Length 25;
 Best Local Similarity 100.0%; Pred. No. 1.1e+03;
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY
 1 GTGCTCCATTGATGC 15
 8 GTGCTCCATTGATGC 22
Db
 8 GTGCTCCATTGATGC 22
Search completed: June 21, 2005, 02:55:37
Job time : 1681 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: June 20, 2005, 20:41:00 ; Search time 434 Seconds
(without alignments)
204.599 Million cell updates/sec

Title: US-10-075-994A-1

Perfect score: 15

Sequence: 1 gtctcctatgatgc 15

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%
Listing first 45 summaries

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2: _geneseq1990s:*

3: _geneseq2000s:*

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5: _geneseq2001bs:*

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11: _geneseq2003ds:*

12: _geneseq2004as:*

13: _geneseq2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	15	100.0	15	2	AAV54043 Human ant
2	15	100.0	15	2	AAV99435 Antisense
3	15	100.0	15	3	AA298661 Human c-r
4	15	100.0	15	6	AA22797 Human c-r
5	15	100.0	15	9	ACC58517 Oligonuc
6	15	100.0	15	9	ADA24233 Human c-r
7	15	100.0	15	10	ADB97458 Sense (AT
8	15	100.0	15	10	ADB97456 Antisense
9	15	100.0	15	10	ADB82830 Immunobcl
10	15	100.0	15	12	ADB90171 Human c-r
11	15	100.0	15	12	AD39690 Oligonuc
12	15	100.0	15	12	AD32025 Antisense
13	15	100.0	15	12	ADP42926 Methylate
14	15	100.0	15	12	AD170154 Oligonuc
15	15	100.0	15	13	ADR88950 Anti c-ra
16	15	100.0	20	2	AA27527 Mouse/rat
17	15	100.0	20	2	AA21157 Mouse and
18	15	100.0	20	3	AA473535 Mouse and
19	15	100.0	20	6	AA444760 Mouse/rat
20	15	100.0	20	10	ADf09751 Mouse/rat

21	15	100.0	20	10	ACD42120 Antisense
22	15	100.0	25	10	ADB97457 Oligo use
23	15	100.0	165	6	ABQ97799 Mouse BS
24	15	100.0	478	9	ACH17231 Human adu
25	15	100.0	597	12	ACH77604 Human gen
26	15	100.0	968	13	ADR60510 Cotton cd
27	15	100.0	1038	10	ADP57946 Human pol
28	15	100.0	2524	10	ADB58074 Toxicity
29	15	100.0	2524	10	ADB52556 Primary r
30	15	100.0	2975	10	AD64080 DNA #1 re
31	15	100.0	2975	12	ADL16251 Raf cDNA.
32	15	100.0	2977	2	AAT30085 Human Raf
33	15	100.0	2977	2	AAT61894 Human Raf
34	15	100.0	2977	2	AAV20439 Human c-r
35	15	100.0	2977	2	AAV99340 Human c-r
36	15	100.0	2977	2	AAV78137 Human c-R
37	15	100.0	2977	3	AAA48654 Human c-r
38	15	100.0	2977	3	AAA73552 Polynucle
39	15	100.0	2977	5	AAV75126 Human c-R
40	15	100.0	2977	6	ABL68999 Pancreas
41	15	100.0	2977	6	ABL57050 Human pro
42	15	100.0	2977	6	ABK72300 Lymphona
43	15	100.0	2977	6	AA168698 Human c-r
44	15	100.0	2977	6	AA444819 Human raf
45	15	100.0	2977	10	ADf18639 Human raf

ALIGNMENTS

RESULT 1	AAV54043	standard; DNA; 15 BP.
ID	AAV54043	
XX	AAV54043;	
AC	02-DEC-1998	(first entry)
DT		
XX		
DE	Human antisense c-raf-1 oligodeoxyribonucleotide.	
XX		
KW	Human; antisense; c-raf-1; oligodeoxyribonucleotide; ODN/oligo;	
KW	Tumour tissue; cancer; radiation therapy; radiosensitive; antisense;	
KW	liposome carrier system; ss.	
OS	Homo sapiens.	
XX		
XX		
FT	Key	Location/Qualifiers
FT	modified_base	1
FT		/*tag= a
FT		/note= "N-terminal base is phosphothioated"
FT	modified_base	15
FT		/*tag= b
FT		/note= "C-terminal base is phosphothioated"
XX		
PN	W09843095-A1.	
XX		
PD	01-OCT-1998.	
XX		
PF	19-MAR-1998;	98WO-US005303.
XX		
PR	21-MAR-1997;	97US-0041192P.
PR	24-OCT-1997;	97US-00957327.
XX		
PI	(GROU) UNIV GEORGETOWN.	
XX		
PI	Kasid U, Gokhale P, Dritschilo A, Rahman A;	
XX	WPI; 1998-532155/45.	
XX		
XX	New cationic liposome composition containing raf oligodeoxynucleotide -	
XX	can be used to directly target tumour tissue and is useful in the	
XX	radiation therapy of cancers.	

CC charges on the particles. One lipid has a (de)protonatable group with Ka
CC much that the lipid is charged at a first pH but neutral at a second pH
CC (particularly near physiological pH) and the buffer maintains this lipid
CC in the charged form (i.e. cationic when the therapeutic agent is anionic
CC in the buffer, or vice versa). The second lipid prevents particle
CC aggregation during formation of the lipid-therapeutic agent particles.
CC The composition is used to introduce therapeutic agents into cells, in
CC vivo or in vitro, particularly to treat or prevent diseases associated
CC with aberrant gene expression in mammals, specifically tumours,
CC inflammation or infection

Query Match	100.0%	Score 15;	DB 2;	Length 15;
Best Local Similarity	100.0%	Pred. No. 1.1e+02;		
Matches 15; Conservative	0;	Mismatches	0;	Indels 0; Gaps 0

Qy	1	GTGCTCCATTGATGC	15
Db	1	GTGCTCCATTGATGC	15

DT 22-MAR-1999 (first entry)

XX	Human c-raf-1 PK therapeutic antisense oligonucleotide sequence ATG-AS
DE	

XX Antisense oligonucleotide; phosphorothioate; inflammatory disease;
KM tumour; gene therapy; aberrant gene expression; treatment;
KM

XX
05 Homo sapiens.

PD 19-NOV-1998.

PR 14-MAY-1997; 97US-00856374.

PA (INEX-) INEX PHARM CORP.

PA (HARA/) HARASYM T.
PA (HOPE/) HOPE M J.

PA (CULL/) CULLIS P R.

PA (SCHE/) SCHERRER P.
PA (SEMP/) SEMPLÉ S. C.
XX

PI Scherrer P, Sempie SC

DR WPI; 2000-225056/20.

XX

XX A method for delivering antisense oligonucleotides to cells using lipid

PT capsules comprising steric barrier lipids.

XX

XX

PS Example 5, Page 57; 99pp; English.

XX

CC This sequence represents an antisense oligonucleotide sequence which has

CC human c-raf-1 protein kinase as its target gene. The oligonucleotide is

CC used in a method for delivering lipid encapsulated therapeutic agents
CC (i.e. antisense oligonucleotides) to mammals. The lipid capsule comprises
CC steric barrier lipids that prevent particle aggregation during lipid
CC nucleic acid formation. The method may be used for the delivery of
CC therapeutic agents to mammalian cells. It is especially suitable for
CC delivering nucleic acid molecules, and in particular antisense molecules
CC which may be administered to down regulate the expression of aberrant
CC genes. The aberrant gene may be ICAM-1, c-myc, c-mycb, raf, erb-B-2,
CC PKC-alpha, IGF-1R, EGFR, VEGF and/or VEG-R-1. The method may be used for
CC the treatment of tumours, inflammatory diseases and/or infectious
CC diseases
XX
SQ Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;
Query Match 100.0%; Score 15; DB 3; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 1 GTGCTCCATTGATGC 15
1 GTGCTCCATTGATGC 15
DB
RESULT 4
AAD22797 standard; DNA, 15 BP.
XX
AC AAD22797;
XX
DT 26-FEB-2002 (first entry)
XX
DE Human c-raf-1 protein kinase antisense oligonucleotide, ATG-AS.
XX
KW Treatment; tumour; lipid-therapeutic agent particle; sphingomyelin;
KM diacylglycerol; phosphatidylcholine; palmitoylcholine; phosphatidylcholine; DSPC;
KW POPC; 1,3-dioleoyl-sn-3-phosphoethanolamine; cholesterol; SM; DOPG;
XX inflammation; c-raf-1 protein kinase gene; human; infectious disease; ss.
XX Homo sapiens.
XX
FH Key location/Qualifiers
FT modified_base 1..20
FT /*tag= a
FT /mod_base= OTHER
FT /note= "Optionally phosphorothioate backbone"
XX
PN US6287591-B1.
XX
PD 11-SEP-2001.
XX
PF 14-MAY-1998; 98US-00078954.
XX
PR 14-MAY-1997; 97US-00856374.
XX
PA (INEX-) INEX PHARM CORP.
XX
PI Semple SC, Klimuk SK, Haraasym T, Hope MD, Ansell SM, Cullis P;
PI Scherrer P, Debeyer D;
XX
DR WPI; 2002-024656/03.
XX
PT Composition useful for treatment of e.g. tumors comprises particles
PT comprising lipid portion and a charged therapeutic agent.
XX
PS Disclosure; Col 15-16; 48pp; English.
XX
CC The invention relates to a composition useful for treatment of e.g.
CC tumours. The composition comprises lipid-therapeutic agent particles
CC comprising a lipid portion and a charged therapeutic agent which is
CC encapsulated in the lipid portion. The lipid portion comprises a first
CC lipid component selected from lipids containing a protonatable or
CC deprotonatable (preferably protonatable) group that has a pKa such that
CC the lipid is in charged form at a first pH and in neutral form at a

CC second pH. The pKa of lipid component is from 4-11. The first lipid
CC component is further selected such that the charged form is cationic when
CC the therapeutic agent is anionic and vice versa; the second lipid
CC component is selected from lipids that prevent particle aggregation
CC during lipid-therapeutic agent particles formation and which exchange out
CC the lipid particle at a rate greater than PEG-CerC20; third lipid
CC component is a neutral lipid selected from diacylglycerol; phosphatidylcholine
CC (DSPC), palmitoylcholine (POPC), 1,2-dioleoyl-sn-3-
CC phosphoethanolamine (DOPG) or SM (sphingomyelin) and a fourth lipid
CC component which is cholesterol. Compositions of the invention are used
CC for treatment or prevention of a disease caused by aberrant expression of
CC a gene preferably ICAM-1 (intracellular adhesion molecule-1), c-myc, c-
CC myb, raf, erb-B-2, PKC-alpha (phosphokinase C-alpha), IGF-1R
CC (insulin growth factor 1-receptor), bcl-2, EGFR (epidermal growth factor
CC receptor), VEGF and VEGF-R-1 (vascular endothelial growth factor receptor
CC 1) in a mammal or by inflammations such as tumour or an infectious
CC disease. The present sequence is an antisense oligonucleotide targeted
CC to human c-raf-1 protein kinase gene
XX
SQ Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;
Query Match 100.0%; Score 15; DB 6; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 1 GTGCTCCATTGATGC 15
1 GTGCTCCATTGATGC 15
DB
RESULT 5
ACCS8517 standard; DNA, 15 BP.
XX
AC ACCS8517;
XX
DT 26-AUG-2003 (first entry)
XX
DE Oligonucleotide ODN #25 (hC-Raf1).
XX
KW Lipid nucleic acid; LNA; mucosal; vaccine; immunostimulant; human;
KM C-Raf-s; ss.
XX
OS Homo sapiens.
XX
FH Key location/Qualifiers
FT modified_base 1..15
FT /*tag= a
FT /mod_base= OTHER
FT /note= "OTHER= optional phosphorothioate nucleotides"
XX
PN WO2003039595-A2.
XX
PD 15-MAY-2003.
XX
PF 07-NOV-2002; 2002WO-CA001717.
XX
PR 07-NOV-2001; 2001US-0337522P.
PR 10-MAY-2002; 2002US-0379343P.
XX
PA (INEX-) INEX PHARM CORP.
XX
PI Semple S, Klimuk S, Yuan Z;
XX
DR WPI; 2003-493235/46.
XX
PT Improved mucosal adjuvant useful in the preparation of vaccine for
PT stimulating an immune response comprises a lipid-nucleic acid formulation
PT containing a nucleic acid component encapsulated by a lipid.
XX
PS Disclosure; Page 21; 71pp; English.
XX
CC The present sequence is that of oligodeoxynucleotide ODN #25 (hC-Raf-1)

```

RESULT 6
ADA24233
ID ADA24233 standard; DNA; 15 BP.
XX
XX ADA24233;
AC
XX
DT 20-NOV-2003 (first entry)
XX
EX Human c-raf-1 protein kinase antisense oligonucleotide SEQ ID NO:16
XX

```

CC	RNA (dsRNA) that includes a reactive group (RG) that can react with a
CC	mobile protein (MP) to form a covalent conjugate or TON/dsRNA and MP; or
CC	(b) TON or dsRNA already conjugated to MP through a covalent bond. Also
CC	described: (1) TON of 15-30 bases that includes (1) a part that binds to
CC	target RNA or DNA and (11) RG; (2) TON of 15-30 bases that includes a
CC	part that binds to target RNA or DNA and is conjugated to MP through a
CC	covalent link; (3) dsRNA that includes RG; and (4) dsRNA that is
CC	conjugated to MP through a covalent link. TON have cytotoxic,
CC	immunosuppressive, virucide, anti-HIV, antibacterial and cardiant
CC	activities. The method is used to treat, or prevent, hyperproliferation
CC	(particularly cancers, solid or haematological), including prevention of
CC	metastatic spread; autoimmune diseases; viral or bacterial infections;
CC	endocrine, neural, cardiovascular, pulmonary or reproductive system
CC	disorders. Also where TON or dsRNA are labelled, they can be used for
CC	diagnosis and monitoring of therapy. When linked to a mobile protein,
CC	TON/dsRNA have better cell entry (via endocytosis or other parts of the
CC	mobile protein metabolic process) and longer therapeutic life, increased
CC	from hours to weeks (the result of increased resistance to nuclease),
CC	without loss of affinity for the target. In many cases immune response to
CC	TON/dsRNA is also reduced, as is non-specific binding to endogenous
CC	proteins. The present sequence represents a human c-rafl-1 antisense
CC	oligonucleotide, which is a specifically claimed TON from the present
CC	invention.
CC	
SQ	Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;
XX	
XX	Query Match 100.0%; Score 15; DB 9; Length 15;
	Best Local Similarity 100.0%; Pred. No. 1,1e+02;
	Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0
OY	1 GTGCTCCATTGATGC 15
D6	1 GTGCTCCATTGATGC 15
RESULT 7	
ID	ADB97458/C
XX	ADB97458 standard; DNA; 15 BP.
AC	
XX	ADB97458;
DE	
DT	04-DEC-2003 (first entry)
XX	
DE	Sense (ATG-S) raf ODN oligodeoxyribonucleotide.
KW	antisense; ATG-S; raf ODN; chemosensitization; tumour tissue;
KW	chemotherapeutic agent; cationic liposome; cationic lipid;
KW	phosphatidylcholine; cholesterol; liposome;
KW	dimethyldioctadecyl ammonium bromide; DDAP;
KW	dimyristoyl trimethyl ammonium propanoate; DMFAP; phosphatidylcholine; PC;
KW	cholesterol; cancer; leukaemia; lymphoma; myeloma; carcinoma; sarcoma;
KW	combination therapy; pre-cancerous lesion; chemotherapy; ss.
OS	Unidentified.
XX	
PN	WO2003070221-A1.
PD	28-AUG-2003.
XX	
PF	14-FEB-2003; 2003WO-US004681.
PR	15-FEB-2002; 2002US-00075994.
XX	
PA	(GEOU) UNIV GEORGETOWN.
PA	(NEOP-) NEOPHARM INC.
P1	Kasid U, Gokhale P, Pei J, Mewani R, Ahmad I, Drischilo A;
P1	Rahman A;
DR	WI; 2003-689738/65.
XX	
PT	Chemosensitization of tumor tissue, useful for treating cancer, e.g.
PT	leukemia, lymphoma or myeloma, comprises administering a chemotherapeutic

PT agent and cationic liposomes containing oligonucleotides.
XX
XX Example 1; Page 18; 77pp; English.
XX
CC The invention relates to a novel method for the chemosensitisation of
CC tumour tissue, comprising administering a chemotherapeutic agent and a
CC composition comprising cationic liposomes consisting of cationic lipid,
CC phosphatidylcholine and cholesterol, where oligonucleotide(s) are
CC encapsulated within the liposome. The invention further relates to a
CC composition comprising liposomes consisting essentially of a cationic
CC lipid like dimethyldioctadecyl ammonium bromide (DDAB) or dimyristoyl
CC trimethyl ammonium propene (DMTAP), phosphatidylcholine (PC),
CC cholesterol, and containing the sequence 5'-GTGCTCCATTGATGC-3', where
CC only the terminal sequences are phosphorothioated. The method is useful
CC for chemosensitisation of a tumour tissue or cancer, including leukaemia,
CC lymphoma, myeloma, carcinoma or sarcoma. The combination therapy may be
CC used for any stage of cancer ranging from pre-cancerous lesions to cancer
CC of advanced stages. This polynucleotide sequence represents the sense
CC (ATG-S) raf ODN oligodeoxyribonucleotide, a cationic liposome of the
CC invention.
XX
SQ Sequence 15 BP; 5 A; 4 C; 4 G; 2 T; 0 U; 0 Other;
XX
Query Match 100.0%; Score 15; DB 10; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 1 GTGCTCCATTGATGC 15
15 GTGCTCCATTGATGC 1
Db
RESULT 8
ADBF7456
ID ADB97456 standard; DNA; 15 BP.
AC ADB97456;
XX
XX 04-DEC-2003 (first entry)
XX
DE Antisense (ATG-AS) raf ODN oligodeoxyribonucleotide.
XX
XX antisense; ATG-AS; raf ODN; chemosensitisation; tumour tissue;
XX chemotherapeutic agent; cationic liposome; cationic lipid;
XX phosphatidylcholine; cholesterol; liposome;
XX dimethyldioctadecyl ammonium bromide; DDAB;
XX dimyristoyl trimethyl ammonium propene; DMTAP; phosphatidylcholine; PC;
XX cholesterol; cancer; leukaemia; lymphoma; myeloma; carcinoma; sarcoma;
XX combination therapy; pre-cancerous lesion; chemotherapy; ss.
XX
XX Unidentified.
XX
XX WO2003070221-A1.
XX
XX 28-AUG-2003.
XX
XX 14-FEB-2003; 2003WO-US004681.
XX
XX 15-FEB-2002; 2002US-00075994.
XX
XX (GBOU) UNIV GEORGETOWN.
XX (NEOP-) NEOPHARM INC.
XX
XX Kasid U, Gokhale P, Pei J, Mewani R, Ahmad I, Drischilo A;
XX Rahman A;
XX
XX WPI; 2003-689738/65.
XX
XX Chemosensitization of tumor tissue, useful for treating cancer, e.g.
XX leukemia, lymphoma or myeloma, comprises administering a chemotherapeutic
XX agent and cationic liposomes containing oligonucleotides.
XX
XX Example 1; Page 18; 77pp; English.

XX
XX The invention relates to a novel method for the chemosensitisation of
XX tumour tissue, comprising administering a chemotherapeutic agent and a
XX composition comprising cationic liposomes consisting of cationic lipid,
XX phosphatidylcholine and cholesterol, where oligonucleotide(s) are
XX encapsulated within the liposome. The invention further relates to a
XX composition comprising liposomes consisting essentially of a cationic
XX lipid like dimethyldioctadecyl ammonium bromide (DDAB) or dimyristoyl
XX trimethyl ammonium propene (DMTAP), phosphatidylcholine (PC),
XX cholesterol, and containing the sequence 5'-GTGCTCCATTGATGC-3', where
XX only the terminal sequences are phosphorothioated. The method is useful
XX for chemosensitisation of a tumour tissue or cancer, including leukaemia,
XX lymphoma, myeloma, carcinoma or sarcoma. The combination therapy may be
XX used for any stage of cancer ranging from pre-cancerous lesions to cancer
XX of advanced stages. This polynucleotide sequence represents the antisense
XX (ATG-AS) raf ODN oligodeoxyribonucleotide, a cationic liposome of the
XX invention.
XX
SQ Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;
XX
Query Match 100.0%; Score 15; DB 10; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 1 GTGCTCCATTGATGC 15
1 GTGCTCCATTGATGC 15
Db
RESULT 9
ADBF82830
ID ADF82830 standard; DNA; 15 BP.
AC ADF82830;
XX
XX 26-FEB-2004 (first entry)
XX
XX Immunostimulant ODN25, component of lipid-nucleic acid vaccine.
XX
XX Immunostimulant; vaccine; lipid-nucleic acid; phosphorothioate; human;
XX C-Raf-S; ss.
XX
XX Synthetic.
XX Homo sapiens.
XX
XX Key location/Qualifiers
XX modified_base 1..15
XX /tag= a
XX /mod_base= OTHER
XX /note= "OTHER= optional phosphorothioate nucleotides"
XX
XX WO2003094829-A2.
XX
XX 20-NOV-2003.
XX
XX 12-MAY-2003; 2003WO-CA000680.
XX
XX 10-MAY-2002; 2002US-0379343P.
XX 07-NOV-2002; 2002US-00290545.
XX 12-MAR-2003; 2003US-0454298P.
XX
XX (INEX-) INEX PHARM CORP.
XX
XX Semple S, Chikh G, Hope MJ, Tam YK;
XX
XX WPI; 2003-903935/82.
XX
XX New pathogen vaccine having a lipid-nucleic acid formulation in
XX combination with at least one microbial antigen, useful for stimulating
XX enhanced responses against bacterial, viral and parasitic infections.
XX
XX Disclosure; SEQ ID NO 25; 138pp; English.
XX

CC The present sequence is that of ODN25 (C-Raf-s) for human C-Raf-s. This
CC is an immunostimulatory oligonucleotide that can be used in lipid-nucleic
CC acid (LNA) vaccines of the invention. Claimed vaccines comprise an LNA
CC formulation in combination with at least one microbial antigen, such as
CC hepatitis B virus surface antigen. The lipid component of the LNA
CC comprises at least one cationic lipid. The oligonucleotide component of
CC the LNA preferably comprises at least one CpG dinucleotide, a methylated
CC cytosine and a phosphorothioate backbone. The vaccine is capable of
CC stimulating Th1 type humoral and cellular immune responses. An enhanced
CC humoral response is demonstrated by a strong early peak of interferon-
CC gamma production observed within hours of vaccine followed by a second
CC stronger peak of interferon-gamma production observed several days later,
CC correlated with antibody isotype switching.

XX Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;

QY Query Match 100.0%; Score 15; DB 10; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 GTGCTCCATTGATGC 15
1 GTGCTCCATTGATGC 15

RESULT 10

ADBE90171 ADE90171 standard; DNA; 15 BP.

AC ADE90171;

DT 12-FEB-2004 (first entry)

DE Human c-raf-1 protein kinase antisense oligonucleotide.

XX ss; lipid-encapsulated therapeutic agent particle;
KW aberrant gene expression; intercellular adhesion molecule; ICAM-1; c-myc;
KW c-myc; ras; raf; erb-B-2; protein kinase C; PKC-alpha;
KW insulin-like growth factor; IGF-IR; epidermal growth factor receptor;
KW EGFR; vascular endothelial growth factor; VEGF; VEGF-R-1; tumour;
KW inflammation; infection; antisense; human.

OS Homo sapiens.

PN US2003129221-A1.

PD 10-JUL-2003.

PF 29-JUN-2001; 2001US-00895480.

PR 14-MAY-1997; 97US-00856374.

PR 14-MAY-1998; 98US-00078954.

XX (SEMP/) SEMPLE S C.
PA (KLIM/) KLIMUK S K.
PA (HARA/) HARASYM T.
PA (HOPE/) HOPE M J.
PA (ANSEL/) ANSELL S M.
PA (CULL/) CULLIS P.
PA (SCHE/) SCHERRER P.
PA (DEBE/) DEBEYER D.

PI Semple SC, Klimuk SK, Harasym T, Hope MJ, Ansell SM, Cullis P;
PI Scherrer P, Debey D;

DR WPI; 2004-031296/03.

XX Preparation of a composition comprising lipid-encapsulated therapeutic
XX agent particles, useful for introducing a nucleic acid into a cell and
XX for treating diseases characterized by aberrant gene expression.

XX Disclosure; SEQ ID NO 15; 52bp; English.

CC The invention relates to a method of preparation of a composition
CC comprising lipid-encapsulated therapeutic agent particles. The
CC composition is useful for introducing a nucleic acid into a cell and for
CC treating diseases characterized by aberrant gene expression (especially
CC intercellular adhesion molecule (ICAM)-1, c-myc, c-myc, ras, raf erb-B-2,
CC protein kinase C (PKC)-alpha, insulin-like growth factor (IGF)-IR,
CC epidermal growth factor receptor (EGFR), vascular endothelial growth
CC factor (VEGF) or VEGF-R-1), e.g. tumours, inflammation or infection. The
CC present sequence represents an antisense oligonucleotide.

XX Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;

QY Query Match 100.0%; Score 15; DB 12; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 GTGCTCCATTGATGC 15
1 GTGCTCCATTGATGC 15

RESULT 11

ADBE39690 ADE39690 standard; DNA; 15 BP.

AC ADE39690;

DT 12-FEB-2004 (first entry)

DE Oligonucleotide ODN 25 (nc-Raf-1) SEQ ID NO:25.

XX cancer; vaccine; lipid-nucleic acid; LNA; tumour-associated antigen;
KW Th-1 based immune response; cytostatic; gene therapy;
KW tumour growth inhibition; tumour; human; ss.

OS Synthetic.

OS Homo sapiens.

PN WO2003094828-A2.

PD 20-NOV-2003.

PF 12-MAY-2003; 2003WO-CA000679.

PR 10-MAY-2002; 2002US-0379343P.

PR 07-NOV-2002; 2002US-00290545.

PR 04-APR-2003; 2003US-0460646P.

XX (INEX-) INEX PHARM CORP.
PA Tam YK, Semple S, Klimuk S, Chikh G;
PA WPI; 2004-011992/01.

PI New cancer vaccine having a lipid-nucleic acid formulation in combination
PI with at least one tumor-associated antigen, useful for stimulating
PI enhanced responses against tumor-associated antigens and for inhibiting
PI tumor growth.

DR WPI; 2004-011992/01.

XX The present invention describes a cancer vaccine (I), which comprises a
XX lipid-nucleic acid (LNA) formulation in combination with at least one
XX tumour-associated antigen that is mixed with or associated with the LNA
XX formulation comprising a lipid component having at least one cationic
XX lipid, and a nucleic acid component comprising at least one
XX oligonucleotide, where the vaccine is capable of stimulating a Th-1 based

PS Example 9; SEQ ID NO 25; 119bp; English.

immune response in vivo to the at least one tumour-associated antigen.
CC (1) has cytostatic activity, and can be used in vaccines, and in gene
CC therapy. The methods and compositions of the present invention can be
CC used for stimulating enhanced responses against tumour-associated
CC antigens and for inhibiting tumour growth. The present sequence
CC represents an oligonucleotide which is used in the exemplification of the
CC present invention.

XX Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 15; DB 12; Length 15;

Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 GTGCTCCATTGATGC 15
|||
1 GTGCTCCATTGATGC 15

RESULT 12

ADF32025 standard; DNA, 15 BP.

XX ADF32025;

DT 26-FEB-2004 (first entry).

DE Antisense oligonucleotide of the invention.

XX platelet; oligonucleotide; Thrombolytic; thrombocytosis; ss.

XX Synthetic.

PN WO200309213-A2.

PD 04-DEC-2003.

PF 19-MAY-2003; 2003WO-US015922.

PR 20-MAY-2002; 2002US-0382411P.

PA (NEOP-) NEOPHARM INC.

XX Gately ST;

XX WPI; 2004-035033/03.

XX Reducing the platelet count in a patient, useful for treating
PT thrombocytosis, comprises administering antisense oligonucleotides
PT inhibiting raf-1 gene with an agent that enhances penetration of the
PT oligonucleotide into cells.

XX Example 1; SEQ ID NO 1; 14pp; English.

XX The present invention relates to reducing the platelet count in a patient
CC comprises preparing a formulation of an oligonucleotide with an agent
CC that enhances penetration of the oligonucleotide into cells, and
CC administering the formulation to a patient having an elevated platelet
CC count. The oligonucleotide is useful for preparing a medication for
CC reducing the platelet count in a patient, particularly for treating
CC thrombocytosis. The present sequence represents an oligonucleotide of the
CC invention.

XX Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 15; DB 12; Length 15;

Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 GTGCTCCATTGATGC 15
|||
1 GTGCTCCATTGATGC 15

RESULT 13

ADF42926 standard; DNA, 15 BP.

XX ADF42926;

DT 11-MAR-2004 (first entry)

DE Methylated immunostimulatory oligonucleotide ODN 25 SEQ ID NO:25.

XX lipid-methylated nucleic acid formulation; immune response;

XX lipid-nucleic acid; vaccine; immunostimulant; cytostatic;

XX antiinflammatory; antiarthritic; gene therapy; cancer; inflammation;

XX arthritis; immunodeficiency disorder; gene therapy; cancer; inflammation;

XX Synthetic.

PN WO2003094963-A2.

PD 20-NOV-2003.

PF 12-MAY-2003; 2003WO-CA000678.

PR 10-MAY-2002; 2002US-0379343P.

PR 07-NOV-2002; 2002US-00290545.

PR 04-APR-2003; 2003US-0460646P.

PA (INEX-) INEX PHARM CORP.

XX Tam YK, Sempke S, Klimuk S, Chikh G;

XX WPI; 2004-142698/14.

XX lipid-methylated nucleic acid formulation for stimulating an immune

XX response in an animal comprises a lipid component and a nucleic acid

XX component comprising a methylated nucleic acid sequence.

XX The present invention describes a lipid-methylated nucleic acid

XX formulation for stimulating an immune response in an animal, comprising a

XX lipid component and a nucleic acid component which is a methylated

XX nucleic acid sequence. Also described: (1) an adjuvant comprising a lipid

XX -nucleic acid (LNA) formulation; (2) a vaccine comprising the LNA

XX formulation in combination with at least one target antigen; (3)

XX stimulating an enhanced host immune response to antigenic stimulation,

XX comprising administering to the host the LNA formulation; (4) stimulating

XX host dendritic cells in vivo, comprising contacting at least one

XX dendritic cell with the lipid-methylated nucleic acid formulation to a

XX host; and (5) simultaneously delivering antigenic and adjuvant immune

XX stimulation to antigen presenting cells, comprising the administration of

XX the LNA formulation associated with a target antigen. The lipid-

XX methylation nucleic acid formulation has immunostimulant, cytostatic,

XX antiinflammatory and antiarthritic activities, and can be used in

XX vaccines, and in gene therapy. The formulation and methods are useful in

XX stimulating a host's immune response to antigenic stimulation, or in

XX activating and/or expanding dendritic cell populations in response to

XX antigenic stimulation. They may be used for treating cancer,

XX inflammation, arthritis or immunodeficiency disorders. The present

XX sequence represents a methylated immunostimulatory oligonucleotide given

XX in the exemplification of the present invention.

XX Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;

Query Match 100.0%; Score 15; DB 12; Length 15;

Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 GTGCTCCATTGATGC 15
|||
1 GTGCTCCATTGATGC 15

RESULT 14
ADL70154
ID ADL70154 standard; DNA; 15 BP.
XX
AC ADL70154;
XX
DT 20-MAY-2004 (first entry)
XX
DE Oligonucleotide antisense to raf.
XX
KM Raf; antisense; liposome; drug delivery; cytostatic; ss.
XX
OS Synthetic.
XX
PN WO2004017944-A1.
XX
PD 04-MAR-2004.
XX
PF 13-AUG-2003; 2003MO-US025293.
XX
PR 23-AUG-2002; 2002US-0405378P.
XX
PA (NEOP-) NEOPHARM INC.
XX
PI Zhang J, Ahmad I;
XX
DR WPI; 2004-257219/24.
XX
PT Treatment of cellular proliferative disease e.g. cancer involves
PT administration of a composition comprising liposomal gemcitabine and
PT negatively charged phospholipid.
XX
PS Disclosure; SEQ ID NO 1; 25pp; English.
XX
CC The present sequence is that of an antisense oligonucleotide to raf. The
CC invention relates to novel gemcitabine compositions and their use in
CC treating proliferative diseases such as cancer, particularly in mammals,
CC especially in humans. The compositions include liposome-entrapped
CC gemcitabine. The cancer is especially lymphoma, ovarian cancer, breast
CC cancer, pancreatic cancer, lung cancer or colon cancer. The liposomal
CC gemcitabine compositions can be used in conjunction with secondary
CC therapeutic agents including antineoplastic, antifungal and antibiotic
CC agents as well as antisense oligonucleotides, especially an antisense
CC oligonucleotide to raf (claimed).
XX
SQ Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;
XX
Query Match 100.0%; Score 15; DB 12; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GTGCTCCATTGATGC 15
|||
Db 1 GTGCTCCATTGATGC 15
|||
RESULT 15
ADR88950
ID ADR88950 standard; DNA; 15 BP.
XX
AC ADR88950;
XX
DT 18-NOV-2004 (first entry)
XX
DE Anti c-raf-1 oligonucleotide.
XX
KM C-raf-1; liposomal; antineoplastic; cytostatic; cancer; antisense; ss.
XX
OS Synthetic.
XX
PN WO2004071466-A2.

XX
PD 26-AUG-2004.
XX
PF 11-FEB-2004; 2004MO-US004555.
XX
PR 11-FEB-2003; 2003US-0446895P.
XX
PA (NEOP-) NEOPHARM INC.
XX
PI Bhamidipati S, Ahmad Z, Ahmad I;
XX
DR WPI; 2004-635030/61.
XX
PT Preparation of liposomal composition used for treating e.g. cancer
PT involves dissolving lipid fraction in water miscible organic solvent and
PT mixing solvent solution with aqueous solution.
XX
PS Disclosure; Page 6; 27pp; English.
XX
CC The invention relates to the preparation of a liposomal composition. The
CC method involves: dissolving a lipid fraction in a water-miscible organic
CC solvent; and mixing the water-miscible organic solvent solution
CC comprising the lipid fraction with an aqueous solution under conditions
CC to form a bulk liposomal composition. The method further involves adding
CC at least one active principal to the water-miscible organic solvent prior
CC to the addition of the lipid fraction, or to the aqueous solution prior,
CC during or after the step (b), size-reducing the bulk liposomal
CC composition to obtain a size-reduced liposomal composition, freeing the
CC liposomal composition of the water-miscible organic solvent by
CC dialfiltration using a tangential flow filtration process and freeze-drying
CC the liposomal preparation. Step (b) involves adding the water-miscible
CC organic solvent solution to the aqueous solution while mixing and mixing
CC or solution following addition of water-miscible solvent comprising the
CC lipid fraction to the aqueous solution while cooling. The active
CC principal comprises at least one antineoplastic or antifungal agent
CC (preferably taxane, camptothecin or their derivatives, especially
CC paclitaxel or docetaxel). The composition is used for the treatment of
CC disease e.g. cancer. The composition eliminates the disease or its
CC symptoms, need not completely eradicate the effects of the disease,
CC reduces the severity of a disease, infection or reduction in the rate by
CC which a disease progresses within a patient. The method permits the
CC production of liposomal formulation on a commercial scale. The present
CC sequence represents an antisense oligonucleotide specific for c-raf-1,
CC that can be used as an active principal.
XX
SQ Sequence 15 BP; 2 A; 4 C; 4 G; 5 T; 0 U; 0 Other;
XX
Query Match 100.0%; Score 15; DB 13; Length 15;
Best Local Similarity 100.0%; Pred. No. 1.1e+02;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GTGCTCCATTGATGC 15
|||
Db 1 GTGCTCCATTGATGC 15
|||
Search completed: June 21, 2005, 02:27:28
Job time : 438 secs